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(21) International Application Number: PCT/IB98/01665		(74) Agent: HALLYBONE, Huw, George; Carpmaels & Ransford, 43 Bloomsbury Square, London WC1A 2RA (GB).	
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(71) Applicant (for all designated States except US): CHIRON S.P.A. [IT/IT]; Via Fiorentina, 1, I-53100 Siena (IT).			
(72) Inventors; and		Published	
(75) Inventors/Applicants (for US only): MASIGNANI, Vega [IT/IT]; Via Pantaneto, 105, I-53100 Siena (IT). RAP-PUOLI, Rino [IT/IT]; Via delle Rocche, 1, Vagliagli, I-53019 Castelnuovo Berardenga (IT). PIZZA, Mariagrazia [IT/IT]; Strada di Montalbuccio, 160, I-53100 Siena (IT). SCARLATO, Vincenzo [IT/IT]; Via Firenze, 3/37, I-53134 Colle Val d'Elsa (IT). GRANDI, Guido [IT/IT]; 9° Strada, 4, I-20090 Segrate (IT).		Without international search report and to be republished upon receipt of that report.	
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(57) Abstract			
<p>The invention provides proteins from <i>Neisseria meningitidis</i> (strains A and B) and from <i>Neisseria gonorrhoeae</i> including amino acid sequences, the corresponding nucleotide sequences, expression data, and serological data. The proteins are useful antigens for vaccines, immunogenic compositions, and/or diagnostics.</p>			

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NEISSERIAL ANTIGENS

This invention relates to antigens from *Neisseria* bacteria.

BACKGROUND ART

Neisseria meningitidis and *Neisseria gonorrhoeae* are non-motile, gram negative diplococci that are pathogenic in humans. *N.meningitidis* colonises the pharynx and causes meningitis (and, occasionally, septicaemia in the absence of meningitis); *N.gonorrhoeae* colonises the genital tract and causes gonorrhea. Although colonising different areas of the body and causing completely different diseases, the two pathogens are closely related, although one feature that clearly differentiates meningococcus from gonococcus is the presence of a polysaccharide capsule that is present in all pathogenic meningococci.

N.gonorrhoeae caused approximately 800,000 cases per year during the period 1983-1990 in the United States alone (chapter by Meitzner & Cohen, "Vaccines Against Gonococcal Infection", In: *New Generation Vaccines*, 2nd edition, ed. Levine, Woodrow, Kaper, & Cobon, Marcel Dekker, New York, 1997, pp.817-842). The disease causes significant morbidity but limited mortality. Vaccination against *N.gonorrhoeae* would be highly desirable, but repeated attempts have failed. The main candidate antigens for this vaccine are surface-exposed proteins such as pili, porins, opacity-associated proteins (Opas) and other surface-exposed proteins such as the Lip, Laz, IgA1 protease and transferrin-binding proteins. The lipooligosaccharide (LOS) has also been suggested as vaccine (Meitzner & Cohen, *supra*).

N.meningitidis causes both endemic and epidemic disease. In the United States the attack rate is 0.6-1 per 100,000 persons per year, and it can be much greater during outbreaks (see Lieberman *et al.* (1996) Safety and Immunogenicity of a Serogroups A/C *Neisseria meningitidis* Oligosaccharide-Protein Conjugate Vaccine in Young Children. *JAMA* 275(19):1499-1503; Schuchat *et al* (1997) Bacterial Meningitis in the United States in 1995. *N Engl J Med* 337(14):970-976). In developing countries, endemic disease rates are much higher and during epidemics incidence rates can reach 500 cases per 100,000 persons per year. Mortality is extremely high, at 10-20% in the United States, and much higher in developing countries. Following the introduction of the conjugate vaccine against *Haemophilus influenzae*, *N. meningitidis* is the major cause of bacterial meningitis at all ages in the United States (Schuchat *et al* (1997) *supra*).

Based on the organism's capsular polysaccharide, 12 serogroups of *N.meningitidis* have been identified. Group A is the pathogen most often implicated in epidemic disease in sub-Saharan Africa. Serogroups B and C are responsible for the vast majority of cases in the United States and in most developed countries. Serogroups W135 and Y are responsible for the rest of the cases in the United States and developed countries. The meningococcal vaccine currently in use is a tetravalent polysaccharide vaccine composed of serogroups A, C, Y and W135. Although efficacious in adolescents and adults, it induces a poor immune response and short duration of protection, and cannot be used in infants [eg. Morbidity and Mortality weekly report, Vol.46, No. RR-5 (1997)]. This is because polysaccharides are T-cell independent antigens that induce a weak immune response that cannot be boosted by repeated immunization. Following the success of the vaccination against *H.influenzae*, conjugate vaccines against serogroups A and C have been developed and are at the final stage of clinical testing (Zollinger WD "New and Improved Vaccines Against Meningococcal Disease" in: *New Generation Vaccines, supra*, pp. 469-488; Lieberman *et al* (1996) *supra*; Costantino *et al* (1992) Development and phase I clinical testing of a conjugate vaccine against meningococcus A and C. *Vaccine* 10:691-698).

Meningococcus B remains a problem, however. This serotype currently is responsible for approximately 50% of total meningitis in the United States, Europe, and South America. The polysaccharide approach cannot be used because the menB capsular polysaccharide is a polymer of $\alpha(2-8)$ -linked *N*-acetyl neuraminic acid that is also present in mammalian tissue. This results in tolerance to the antigen; indeed, if an immune response were elicited, it would be anti-self, and therefore undesirable. In order to avoid induction of autoimmunity and to induce a protective immune response, the capsular polysaccharide has, for instance, been chemically modified substituting the *N*-acetyl groups with *N*-propionyl groups, leaving the specific antigenicity unaltered (Romero & Outschoorn (1994) Current status of Meningococcal group B vaccine candidates: capsular or non-capsular? *Clin Microbiol Rev* 7(4):559-575).

Alternative approaches to menB vaccines have used complex mixtures of outer membrane proteins (OMPs), containing either the OMPs alone, or OMPs enriched in porins, or deleted of the class 4 OMPs that are believed to induce antibodies that block bactericidal activity. This approach produces vaccines that are not well characterized. They are able to protect against the homologous strain, but are not effective at large where there are many antigenic variants of the outer membrane proteins. To overcome the antigenic variability, multivalent vaccines containing up to nine different

porins have been constructed (eg. Poolman JT (1992) Development of a meningococcal vaccine. *Infect. Agents Dis.* 4:13-28). Additional proteins to be used in outer membrane vaccines have been the opa and opc proteins, but none of these approaches have been able to overcome the antigenic variability (eg. Ala'Aldeen & Borriello (1996) The meningococcal transferrin-binding proteins 1 and 2 are both surface exposed and generate bactericidal antibodies capable of killing homologous and heterologous strains. *Vaccine* 14(1):49-53).

A certain amount of sequence data is available for meningococcal and gonococcal genes and proteins (eg. EP-A-0467714, WO96/29412), but this is by no means complete. The provision of further sequences could provide an opportunity to identify secreted or surface-exposed proteins that are presumed targets for the immune system and which are not antigenically variable. For instance, some of the identified proteins could be components of efficacious vaccines against meningococcus B, some could be components of vaccines against all meningococcal serotypes, and others could be components of vaccines against all pathogenic *Neisseriae*.

THE INVENTION

The invention provides proteins comprising the Neisserial amino acid sequences disclosed in the examples. These sequences relate to *N.meningitidis* or *N.gonorrhoeae*.

It also provides proteins comprising sequences homologous (*ie.* having sequence identity) to the Neisserial amino acid sequences disclosed in the examples. Depending on the particular sequence, the degree of identity is preferably greater than 50% (eg. 65%, 80%, 90%, or more). These homologous proteins include mutants and allelic variants of the sequences disclosed in the examples. Typically, 50% identity or more between two proteins is considered to be an indication of functional equivalence. Identity between the proteins is preferably determined by the Smith-Waterman homology search algorithm as implemented in the MPSRCH program (Oxford Molecular), using an affine gap search with parameters *gap open penalty*=12 and *gap extension penalty*=1.

The invention further provides proteins comprising fragments of the Neisserial amino acid sequences disclosed in the examples. The fragments should comprise at least *n* consecutive amino acids from the sequences and, depending on the particular sequence, *n* is 7 or more (eg. 8, 10, 12, 14, 16, 18, 20 or more). Preferably the fragments comprise an epitope from the sequence.

The proteins of the invention can, of course, be prepared by various means (*eg.* recombinant expression, purification from cell culture, chemical synthesis *etc.*) and in various forms (*eg.* native, fusions *etc.*). They are preferably prepared in substantially pure or isolated form (*ie.* substantially free from other Neisserial or host cell proteins)

- 5 According to a further aspect, the invention provides antibodies which bind to these proteins. These may be polyclonal or monoclonal and may be produced by any suitable means.

According to a further aspect, the invention provides nucleic acid comprising the Neisserial nucleotide sequences disclosed in the examples. In addition, the invention provides nucleic acid comprising sequences homologous (*ie.* having sequence identity) to the Neisserial nucleotide
10 sequences disclosed in the examples.

Furthermore, the invention provides nucleic acid which can hybridise to the Neisserial nucleic acid disclosed in the examples, preferably under "high stringency" conditions (*eg.* 65°C in a 0.1xSSC, 0.5% SDS solution).

Nucleic acid comprising fragments of these sequences are also provided. These should comprise
15 at least n consecutive nucleotides from the Neisserial sequences and, depending on the particular sequence, n is 10 or more (*eg.* 12, 14, 15, 18, 20, 25, 30, 35, 40 or more).

According to a further aspect, the invention provides nucleic acid encoding the proteins and protein fragments of the invention.

It should also be appreciated that the invention provides nucleic acid comprising sequences
20 complementary to those described above (*eg.* for antisense or probing purposes).

Nucleic acid according to the invention can, of course, be prepared in many ways (*eg.* by chemical synthesis, from genomic or cDNA libraries, from the organism itself *etc.*) and can take various forms (*eg.* single stranded, double stranded, vectors, probes *etc.*).

In addition, the term "nucleic acid" includes DNA and RNA, and also their analogues, such as
25 those containing modified backbones, and also peptide nucleic acids (PNA) *etc.*

According to a further aspect, the invention provides vectors comprising nucleotide sequences of the invention (eg. expression vectors) and host cells transformed with such vectors.

According to a further aspect, the invention provides compositions comprising protein, antibody, and/or nucleic acid according to the invention. These compositions may be suitable as vaccines,
5 for instance, or as diagnostic reagents, or as immunogenic compositions.

The invention also provides nucleic acid, protein, or antibody according to the invention for use as medicaments (eg. as vaccines) or as diagnostic reagents. It also provides the use of nucleic acid, protein, or antibody according to the invention in the manufacture of: (i) a medicament for treating or preventing infection due to Neisserial bacteria; (ii) a diagnostic reagent for detecting the
10 presence of Neisserial bacteria or of antibodies raised against Neisserial bacteria; and/or (iii) a reagent which can raise antibodies against Neisserial bacteria. Said Neisserial bacteria may be any species or strain (such as *N.gonorrhoeae*, or any strain of *N.meningitidis*, such as strain A, strain B or strain C).

The invention also provides a method of treating a patient, comprising administering to the patient
15 a therapeutically effective amount of nucleic acid, protein, and/or antibody according to the invention.

According to further aspects, the invention provides various processes.

A process for producing proteins of the invention is provided, comprising the step of culturing a host cell according to the invention under conditions which induce protein expression.

20 A process for producing protein or nucleic acid of the invention is provided, wherein the the protein or nucleic acid is synthesised in part or in whole using chemical means.

A process for detecting polynucleotides of the invention is provided, comprising the steps of: (a) contacting a nucleic probe according to the invention with a biological sample under hybridizing conditions to form duplexes; and (b) detecting said duplexes.

25 A process for detecting proteins of the invention is provided, comprising the steps of: (a) contacting an antibody according to the invention with a biological sample under conditions suitable for the formation of an antibody-antigen complexes; and (b) detecting said complexes.

A summary of standard techniques and procedures which may be employed in order to perform the invention (eg. to utilise the disclosed sequences for vaccination or diagnostic purposes) follows. This summary is not a limitation on the invention but, rather, gives examples that may be used, but are not required.

5 General

The practice of the present invention will employ, unless otherwise indicated, conventional techniques of molecular biology, microbiology, recombinant DNA, and immunology, which are within the skill of the art. Such techniques are explained fully in the literature eg. Sambrook *Molecular Cloning; A Laboratory Manual, Second Edition* (1989); *DNA Cloning, Volumes I and*
10 *ii* (D.N Glover ed. 1985); *Oligonucleotide Synthesis* (M.J. Gait ed, 1984); *Nucleic Acid Hybridization* (B.D. Hames & S.J. Higgins eds. 1984); *Transcription and Translation* (B.D. Hames & S.J. Higgins eds. 1984); *Animal Cell Culture* (R.I. Freshney ed. 1986); *Immobilized Cells and Enzymes* (IRL Press, 1986); B. Perbal, *A Practical Guide to Molecular Cloning* (1984); the *Methods in Enzymology* series (Academic Press, Inc.), especially volumes 154 & 155; *Gene*
15 *Transfer Vectors for Mammalian Cells* (J.H. Miller and M.P. Calos eds. 1987, Cold Spring Harbor Laboratory); Mayer and Walker, eds. (1987), *Immunochemical Methods in Cell and Molecular Biology* (Academic Press, London); Scopes, (1987) *Protein Purification: Principles and Practice*, Second Edition (Springer-Verlag, N.Y.), and *Handbook of Experimental Immunology, Volumes I-IV* (D.M. Weir and C. C. Blackwell eds 1986).

20 Standard abbreviations for nucleotides and amino acids are used in this specification.

All publications, patents, and patent applications cited herein are incorporated in full by reference. In particular, the contents of UK patent applications 9723516.2, 9724190.5, 9724386.9, 9725158.1, 9726147.3, 9800759.4, and 9819016.8 are incorporated herein.

Definitions

25 A composition containing X is "substantially free of" Y when at least 85% by weight of the total X+Y in the composition is X. Preferably, X comprises at least about 90% by weight of the total of X+Y in the composition, more preferably at least about 95% or even 99% by weight.

The term "comprising" means "including" as well as "consisting" eg. a composition "comprising" X may consist exclusively of X or may include something additional to X, such as X+Y.

The term "heterologous" refers to two biological components that are not found together in nature. The components may be host cells, genes, or regulatory regions, such as promoters. Although the heterologous components are not found together in nature, they can function together, as when a promoter heterologous to a gene is operably linked to the gene. Another example is where a
5 Neisserial sequence is heterologous to a mouse host cell. A further examples would be two epitopes from the same or different proteins which have been assembled in a single protein in an arrangement not found in nature.

An "origin of replication" is a polynucleotide sequence that initiates and regulates replication of polynucleotides, such as an expression vector. The origin of replication behaves as an autonomous
10 unit of polynucleotide replication within a cell, capable of replication under its own control. An origin of replication may be needed for a vector to replicate in a particular host cell. With certain origins of replication, an expression vector can be reproduced at a high copy number in the presence of the appropriate proteins within the cell. Examples of origins are the autonomously replicating sequences, which are effective in yeast; and the viral T-antigen, effective in COS-7
15 cells.

A "mutant" sequence is defined as DNA, RNA or amino acid sequence differing from but having sequence identity with the native or disclosed sequence. Depending on the particular sequence, the degree of sequence identity between the native or disclosed sequence and the mutant sequence is preferably greater than 50% (eg. 60%, 70%, 80%, 90%, 95%, 99% or more, calculated using the
20 Smith-Waterman algorithm as described above). As used herein, an "allelic variant" of a nucleic acid molecule, or region, for which nucleic acid sequence is provided herein is a nucleic acid molecule, or region, that occurs essentially at the same locus in the genome of another or second isolate, and that, due to natural variation caused by, for example, mutation or recombination, has a similar but not identical nucleic acid sequence. A coding region allelic variant typically encodes
25 a protein having similar activity to that of the protein encoded by the gene to which it is being compared. An allelic variant can also comprise an alteration in the 5' or 3' untranslated regions of the gene, such as in regulatory control regions (eg. see US patent 5,753,235).

Expression systems

The Neisserial nucleotide sequences can be expressed in a variety of different expression systems;
30 for example those used with mammalian cells, baculoviruses, plants, bacteria, and yeast.

i. Mammalian Systems

Mammalian expression systems are known in the art. A mammalian promoter is any DNA sequence capable of binding mammalian RNA polymerase and initiating the downstream (3') transcription of a coding sequence (eg. structural gene) into mRNA. A promoter will have a transcription initiating region, which is usually placed proximal to the 5' end of the coding sequence, and a TATA box, usually located 25-30 base pairs (bp) upstream of the transcription initiation site. The TATA box is thought to direct RNA polymerase II to begin RNA synthesis at the correct site. A mammalian promoter will also contain an upstream promoter element, usually located within 100 to 200 bp upstream of the TATA box. An upstream promoter element determines the rate at which transcription is initiated and can act in either orientation [Sambrook et al. (1989) "Expression of Cloned Genes in Mammalian Cells." In *Molecular Cloning: A Laboratory Manual*, 2nd ed.].

Mammalian viral genes are often highly expressed and have a broad host range; therefore sequences encoding mammalian viral genes provide particularly useful promoter sequences. Examples include the SV40 early promoter, mouse mammary tumor virus LTR promoter, adenovirus major late promoter (Ad MLP), and herpes simplex virus promoter. In addition, sequences derived from non-viral genes, such as the murine metallothionein gene, also provide useful promoter sequences. Expression may be either constitutive or regulated (inducible), depending on the promoter can be induced with glucocorticoid in hormone-responsive cells.

The presence of an enhancer element (enhancer), combined with the promoter elements described above, will usually increase expression levels. An enhancer is a regulatory DNA sequence that can stimulate transcription up to 1000-fold when linked to homologous or heterologous promoters, with synthesis beginning at the normal RNA start site. Enhancers are also active when they are placed upstream or downstream from the transcription initiation site, in either normal or flipped orientation, or at a distance of more than 1000 nucleotides from the promoter [Maniatis et al. (1987) *Science* 236:1237; Alberts et al. (1989) *Molecular Biology of the Cell*, 2nd ed.]. Enhancer elements derived from viruses may be particularly useful, because they usually have a broader host range. Examples include the SV40 early gene enhancer [Dijkema et al (1985) *EMBO J.* 4:761] and the enhancer/promoters derived from the long terminal repeat (LTR) of the Rous Sarcoma Virus [Gorman et al. (1982b) *Proc. Natl. Acad. Sci.* 79:6777] and from human cytomegalovirus [Boshart et al. (1985) *Cell* 41:521]. Additionally, some enhancers are regulatable and become active only

in the presence of an inducer, such as a hormone or metal ion [Sassone-Corsi and Borelli (1986) *Trends Genet.* 2:215; Maniatis et al. (1987) *Science* 236:1237].

A DNA molecule may be expressed intracellularly in mammalian cells. A promoter sequence may be directly linked with the DNA molecule, in which case the first amino acid at the N-terminus of the recombinant protein will always be a methionine, which is encoded by the ATG start codon. If desired, the N-terminus may be cleaved from the protein by *in vitro* incubation with cyanogen bromide.

Alternatively, foreign proteins can also be secreted from the cell into the growth media by creating chimeric DNA molecules that encode a fusion protein comprised of a leader sequence fragment that provides for secretion of the foreign protein in mammalian cells. Preferably, there are processing sites encoded between the leader fragment and the foreign gene that can be cleaved either *in vivo* or *in vitro*. The leader sequence fragment usually encodes a signal peptide comprised of hydrophobic amino acids which direct the secretion of the protein from the cell. The adenovirus tripartite leader is an example of a leader sequence that provides for secretion of a foreign protein in mammalian cells.

Usually, transcription termination and polyadenylation sequences recognized by mammalian cells are regulatory regions located 3' to the translation stop codon and thus, together with the promoter elements, flank the coding sequence. The 3' terminus of the mature mRNA is formed by site-specific post-transcriptional cleavage and polyadenylation [Birnstiel et al. (1985) *Cell* 41:349; Proudfoot and Whitelaw (1988) "Termination and 3' end processing of eukaryotic RNA. In *Transcription and splicing* (ed. B.D. Hames and D.M. Glover); Proudfoot (1989) *Trends Biochem. Sci.* 14:105]. These sequences direct the transcription of an mRNA which can be translated into the polypeptide encoded by the DNA. Examples of transcription terminator/polyadenylation signals include those derived from SV40 [Sambrook et al (1989) "Expression of cloned genes in cultured mammalian cells." In *Molecular Cloning: A Laboratory Manual*].

Usually, the above described components, comprising a promoter, polyadenylation signal, and transcription termination sequence are put together into expression constructs. Enhancers, introns with functional splice donor and acceptor sites, and leader sequences may also be included in an expression construct, if desired. Expression constructs are often maintained in a replicon, such as an extrachromosomal element (eg. plasmids) capable of stable maintenance in a host, such as mammalian cells or bacteria. Mammalian replication systems include those derived from animal

viruses, which require trans-acting factors to replicate. For example, plasmids containing the replication systems of papovaviruses, such as SV40 [Gluzman (1981) *Cell* 23:175] or polyomavirus, replicate to extremely high copy number in the presence of the appropriate viral T antigen. Additional examples of mammalian replicons include those derived from bovine papillomavirus and Epstein-Barr virus. Additionally, the replicon may have two replicaton systems, thus allowing it to be maintained, for example, in mammalian cells for expression and in a prokaryotic host for cloning and amplification. Examples of such mammalian-bacteria shuttle vectors include pMT2 [Kaufman et al. (1989) *Mol. Cell. Biol.* 9:946] and pHEBO [Shimizu et al. (1986) *Mol. Cell. Biol.* 6:1074].

10 The transformation procedure used depends upon the host to be transformed. Methods for introduction of heterologous polynucleotides into mammalian cells are known in the art and include dextran-mediated transfection, calcium phosphate precipitation, polybrene mediated transfection, protoplast fusion, electroporation, encapsulation of the polynucleotide(s) in liposomes, and direct microinjection of the DNA into nuclei.

15 Mammalian cell lines available as hosts for expression are known in the art and include many immortalized cell lines available from the American Type Culture Collection (ATCC), including but not limited to, Chinese hamster ovary (CHO) cells, HeLa cells, baby hamster kidney (BHK) cells, monkey kidney cells (COS), human hepatocellular carcinoma cells (*eg.* Hep G2), and a number of other cell lines.

20 ii. Baculovirus Systems

The polynucleotide encoding the protein can also be inserted into a suitable insect expression vector, and is operably linked to the control elements within that vector. Vector construction employs techniques which are known in the art. Generally, the components of the expression system include a transfer vector, usually a bacterial plasmid, which contains both a fragment of the baculovirus genome, and a convenient restriction site for insertion of the heterologous gene or genes to be expressed; a wild type baculovirus with a sequence homologous to the baculovirus-specific fragment in the transfer vector (this allows for the homologous recombination of the heterologous gene in to the baculovirus genome); and appropriate insect host cells and growth media.

After inserting the DNA sequence encoding the protein into the transfer vector, the vector and the wild type viral genome are transfected into an insect host cell where the vector and viral genome are allowed to recombine. The packaged recombinant virus is expressed and recombinant plaques

are identified and purified. Materials and methods for baculovirus/insect cell expression systems are commercially available in kit form from, *inter alia*, Invitrogen, San Diego CA ("MaxBac" kit). These techniques are generally known to those skilled in the art and fully described in Summers and Smith, *Texas Agricultural Experiment Station Bulletin No. 1555* (1987) (hereinafter "Summers and Smith").

Prior to inserting the DNA sequence encoding the protein into the baculovirus genome, the above described components, comprising a promoter, leader (if desired), coding sequence of interest, and transcription termination sequence, are usually assembled into an intermediate transplacement construct (transfer vector). This construct may contain a single gene and operably linked regulatory elements; multiple genes, each with its own set of operably linked regulatory elements; or multiple genes, regulated by the same set of regulatory elements. Intermediate transplacement constructs are often maintained in a replicon, such as an extrachromosomal element (*eg.* plasmids) capable of stable maintenance in a host, such as a bacterium. The replicon will have a replication system, thus allowing it to be maintained in a suitable host for cloning and amplification.

Currently, the most commonly used transfer vector for introducing foreign genes into AcNPV is pAc373. Many other vectors, known to those of skill in the art, have also been designed. These include, for example, pVL985 (which alters the polyhedrin start codon from ATG to ATT, and which introduces a BamHI cloning site 32 basepairs downstream from the ATT; see Luckow and Summers, *Virology* (1989) 17:31.

The plasmid usually also contains the polyhedrin polyadenylation signal (Miller et al. (1988) *Ann. Rev. Microbiol.*, 42:177) and a prokaryotic ampicillin-resistance (*amp*) gene and origin of replication for selection and propagation in *E. coli*.

Baculovirus transfer vectors usually contain a baculovirus promoter. A baculovirus promoter is any DNA sequence capable of binding a baculovirus RNA polymerase and initiating the downstream (5' to 3') transcription of a coding sequence (*eg.* structural gene) into mRNA. A promoter will have a transcription initiation region which is usually placed proximal to the 5' end of the coding sequence. This transcription initiation region usually includes an RNA polymerase binding site and a transcription initiation site. A baculovirus transfer vector may also have a second domain called an enhancer, which, if present, is usually distal to the structural gene. Expression may be either regulated or constitutive.

Structural genes, abundantly transcribed at late times in a viral infection cycle, provide particularly useful promoter sequences. Examples include sequences derived from the gene encoding the viral polyhedron protein, Friesen et al., (1986) "The Regulation of Baculovirus Gene Expression," in: *The Molecular Biology of Baculoviruses* (ed. Walter Doerfler); EPO Publ. Nos. 127 839 and 155
5 476; and the gene encoding the p10 protein, Vlaskovits et al., (1988), *J. Gen. Virol.* 69:765.

DNA encoding suitable signal sequences can be derived from genes for secreted insect or baculovirus proteins, such as the baculovirus polyhedrin gene (Carbonell et al. (1988) *Gene*, 73:409). Alternatively, since the signals for mammalian cell posttranslational modifications (such as signal peptide cleavage, proteolytic cleavage, and phosphorylation) appear to be recognized by
10 insect cells, and the signals required for secretion and nuclear accumulation also appear to be conserved between the invertebrate cells and vertebrate cells, leaders of non-insect origin, such as those derived from genes encoding human α -interferon, Maeda et al., (1985), *Nature* 315:592; human gastrin-releasing peptide, Lebacqz-Verheyden et al., (1988), *Molec. Cell. Biol.* 8:3129; human IL-2, Smith et al., (1985) *Proc. Nat'l Acad. Sci. USA*, 82:8404; mouse IL-3, (Miyajima et al., (1987) *Gene* 58:273; and human glucocerebrosidase, Martin et al. (1988) *DNA*, 7:99, can also
15 be used to provide for secretion in insects.

A recombinant polypeptide or polyprotein may be expressed intracellularly or, if it is expressed with the proper regulatory sequences, it can be secreted. Good intracellular expression of nonfused foreign proteins usually requires heterologous genes that ideally have a short leader sequence
20 containing suitable translation initiation signals preceding an ATG start signal. If desired, methionine at the N-terminus may be cleaved from the mature protein by *in vitro* incubation with cyanogen bromide.

Alternatively, recombinant polyproteins or proteins which are not naturally secreted can be secreted from the insect cell by creating chimeric DNA molecules that encode a fusion protein comprised
25 of a leader sequence fragment that provides for secretion of the foreign protein in insects. The leader sequence fragment usually encodes a signal peptide comprised of hydrophobic amino acids which direct the translocation of the protein into the endoplasmic reticulum.

After insertion of the DNA sequence and/or the gene encoding the expression product precursor of the protein, an insect cell host is co-transformed with the heterologous DNA of the transfer
30 vector and the genomic DNA of wild type baculovirus -- usually by co-transfection. The promoter

- and transcription termination sequence of the construct will usually comprise a 2-5kb section of the baculovirus genome. Methods for introducing heterologous DNA into the desired site in the baculovirus virus are known in the art. (See Summers and Smith *supra*; Ju et al. (1987); Smith et al., *Mol. Cell. Biol.* (1983) 3:2156; and Luckow and Summers (1989)). For example, the insertion
- 5 can be into a gene such as the polyhedrin gene, by homologous double crossover recombination; insertion can also be into a restriction enzyme site engineered into the desired baculovirus gene. Miller et al., (1989), *Bioessays* 4:91. The DNA sequence, when cloned in place of the polyhedrin gene in the expression vector, is flanked both 5' and 3' by polyhedrin-specific sequences and is positioned downstream of the polyhedrin promoter.
- 10 The newly formed baculovirus expression vector is subsequently packaged into an infectious recombinant baculovirus. Homologous recombination occurs at low frequency (between about 1% and about 5%); thus, the majority of the virus produced after cotransfection is still wild-type virus. Therefore, a method is necessary to identify recombinant viruses. An advantage of the expression system is a visual screen allowing recombinant viruses to be distinguished. The polyhedrin protein,
- 15 which is produced by the native virus, is produced at very high levels in the nuclei of infected cells at late times after viral infection. Accumulated polyhedrin protein forms occlusion bodies that also contain embedded particles. These occlusion bodies, up to 15 μ m in size, are highly refractile, giving them a bright shiny appearance that is readily visualized under the light microscope. Cells infected with recombinant viruses lack occlusion bodies. To distinguish recombinant virus from
- 20 wild-type virus, the transfection supernatant is plaqued onto a monolayer of insect cells by techniques known to those skilled in the art. Namely, the plaques are screened under the light microscope for the presence (indicative of wild-type virus) or absence (indicative of recombinant virus) of occlusion bodies. "Current Protocols in Microbiology" Vol. 2 (Ausubel et al. eds) at 16.8 (Supp. 10, 1990); Summers and Smith, *supra*; Miller et al. (1989).
- 25 Recombinant baculovirus expression vectors have been developed for infection into several insect cells. For example, recombinant baculoviruses have been developed for, *inter alia*: *Aedes aegypti*, *Autographa californica*, *Bombyx mori*, *Drosophila melanogaster*, *Spodoptera frugiperda*, and *Trichoplusia ni* (WO 89/046699; Carbonell et al., (1985) *J. Virol.* 56:153; Wright (1986) *Nature* 321:718; Smith et al., (1983) *Mol. Cell. Biol.* 3:2156; and see generally, Fraser, *et al.* (1989) *In Vitro Cell. Dev. Biol.* 25:225).
- 30

Cells and cell culture media are commercially available for both direct and fusion expression of heterologous polypeptides in a baculovirus/expression system; cell culture technology is generally known to those skilled in the art. *See, eg. Summers and Smith supra.*

The modified insect cells may then be grown in an appropriate nutrient medium, which allows for
5 stable maintenance of the plasmid(s) present in the modified insect host. Where the expression product gene is under inducible control, the host may be grown to high density, and expression induced. Alternatively, where expression is constitutive, the product will be continuously expressed into the medium and the nutrient medium must be continuously circulated, while removing the product of interest and augmenting depleted nutrients. The product may be purified by such techniques as
10 chromatography, eg. HPLC, affinity chromatography, ion exchange chromatography, etc.; electrophoresis; density gradient centrifugation; solvent extraction, or the like. As appropriate, the product may be further purified, as required, so as to remove substantially any insect proteins which are also secreted in the medium or result from lysis of insect cells, so as to provide a product which is at least substantially free of host debris, eg. proteins, lipids and polysaccharides.

15 In order to obtain protein expression, recombinant host cells derived from the transformants are incubated under conditions which allow expression of the recombinant protein encoding sequence. These conditions will vary, dependent upon the host cell selected. However, the conditions are readily ascertainable to those of ordinary skill in the art, based upon what is known in the art.

iii. Plant Systems

20 There are many plant cell culture and whole plant genetic expression systems known in the art. Exemplary plant cellular genetic expression systems include those described in patents, such as: US 5,693,506; US 5,659,122; and US 5,608,143. Additional examples of genetic expression in plant cell culture has been described by Zenk, *Phytochemistry* 30:3861-3863 (1991). Descriptions of plant protein signal peptides may be found in addition to the references described above in
25 Vaulcombe et al., *Mol. Gen. Genet.* 209:33-40 (1987); Chandler et al., *Plant Molecular Biology* 3:407-418 (1984); Rogers, *J. Biol. Chem.* 260:3731-3738 (1985); Rothstein et al., *Gene* 55:353-356 (1987); Whittier et al., *Nucleic Acids Research* 15:2515-2535 (1987); Wirsal et al., *Molecular Microbiology* 3:3-14 (1989); Yu et al., *Gene* 122:247-253 (1992). A description of the regulation of plant gene expression by the phytohormone, gibberellic acid and secreted enzymes induced by
30 gibberellic acid can be found in R.L. Jones and J. MacMillan, *Gibberellins: in: Advanced Plant Physiology*,. Malcolm B. Wilkins, ed., 1984 Pitman Publishing Limited, London, pp. 21-52.

References that describe other metabolically-regulated genes: Sheen, *Plant Cell*, 2:1027-1038(1990); Maas et al., *EMBO J.* 9:3447-3452 (1990); Benkel and Hickey, *Proc. Natl. Acad. Sci.* 84:1337-1339 (1987)

Typically, using techniques known in the art, a desired polynucleotide sequence is inserted into an expression cassette comprising genetic regulatory elements designed for operation in plants. The
5 expression cassette is inserted into a desired expression vector with companion sequences upstream and downstream from the expression cassette suitable for expression in a plant host. The companion sequences will be of plasmid or viral origin and provide necessary characteristics to the vector to permit the vectors to move DNA from an original cloning host, such as bacteria, to the
10 desired plant host. The basic bacterial/plant vector construct will preferably provide a broad host range prokaryote replication origin; a prokaryote selectable marker; and, for *Agrobacterium* transformations, T DNA sequences for *Agrobacterium*-mediated transfer to plant chromosomes. Where the heterologous gene is not readily amenable to detection, the construct will preferably also have a selectable marker gene suitable for determining if a plant cell has been transformed. A
15 general review of suitable markers, for example for the members of the grass family, is found in Wilmink and Dons, 1993, *Plant Mol. Biol. Repr.*, 11(2):165-185.

Sequences suitable for permitting integration of the heterologous sequence into the plant genome are also recommended. These might include transposon sequences and the like for homologous recombination as well as Ti sequences which permit random insertion of a heterologous expression
20 cassette into a plant genome. Suitable prokaryote selectable markers include resistance toward antibiotics such as ampicillin or tetracycline. Other DNA sequences encoding additional functions may also be present in the vector, as is known in the art.

The nucleic acid molecules of the subject invention may be included into an expression cassette for expression of the protein(s) of interest. Usually, there will be only one expression cassette,
25 although two or more are feasible. The recombinant expression cassette will contain in addition to the heterologous protein encoding sequence the following elements, a promoter region, plant 5' untranslated sequences, initiation codon depending upon whether or not the structural gene comes equipped with one, and a transcription and translation termination sequence. Unique restriction enzyme sites at the 5' and 3' ends of the cassette allow for easy insertion into a pre-existing vector.

A heterologous coding sequence may be for any protein relating to the present invention. The sequence encoding the protein of interest will encode a signal peptide which allows processing and translocation of the protein, as appropriate, and will usually lack any sequence which might result in the binding of the desired protein of the invention to a membrane. Since, for the most part, the transcriptional initiation region will be for a gene which is expressed and translocated during germination, by employing the signal peptide which provides for translocation, one may also provide for translocation of the protein of interest. In this way, the protein(s) of interest will be translocated from the cells in which they are expressed and may be efficiently harvested. Typically secretion in seeds are across the aleurone or scutellar epithelium layer into the endosperm of the seed. While it is not required that the protein be secreted from the cells in which the protein is produced, this facilitates the isolation and purification of the recombinant protein.

Since the ultimate expression of the desired gene product will be in a eucaryotic cell it is desirable to determine whether any portion of the cloned gene contains sequences which will be processed out as introns by the host's splicosome machinery. If so, site-directed mutagenesis of the "intron" region may be conducted to prevent losing a portion of the genetic message as a false intron code, Reed and Maniatis, *Cell* 41:95-105, 1985.

The vector can be microinjected directly into plant cells by use of micropipettes to mechanically transfer the recombinant DNA. Crossway, *Mol. Gen. Genet.*, 202:179-185, 1985. The genetic material may also be transferred into the plant cell by using polyethylene glycol, Krens, et al., *Nature*, 296, 72-74, 1982. Another method of introduction of nucleic acid segments is high velocity ballistic penetration by small particles with the nucleic acid either within the matrix of small beads or particles, or on the surface, Klein, et al., *Nature*, 327, 70-73, 1987 and Knudsen and Muller, 1991, *Planta*, 185:330-336 teaching particle bombardment of barley endosperm to create transgenic barley. Yet another method of introduction would be fusion of protoplasts with other entities, either minicells, cells, lysosomes or other fusible lipid-surfaced bodies, Fraley, et al., *Proc. Natl. Acad. Sci. USA*, 79, 1859-1863, 1982.

The vector may also be introduced into the plant cells by electroporation. (Fromm et al., *Proc. Natl. Acad. Sci. USA* 82:5824, 1985). In this technique, plant protoplasts are electroporated in the presence of plasmids containing the gene construct. Electrical impulses of high field strength reversibly permeabilize biomembranes allowing the introduction of the plasmids. Electroporated plant protoplasts reform the cell wall, divide, and form plant callus.

All plants from which protoplasts can be isolated and cultured to give whole regenerated plants can be transformed by the present invention so that whole plants are recovered which contain the transferred gene. It is known that practically all plants can be regenerated from cultured cells or tissues, including but not limited to all major species of sugarcane, sugar beet, cotton, fruit and
5 other trees, legumes and vegetables. Some suitable plants include, for example, species from the genera *Fragaria*, *Lotus*, *Medicago*, *Onobrychis*, *Trifolium*, *Trigonella*, *Vigna*, *Citrus*, *Linum*, *Geranium*, *Manihot*, *Daucus*, *Arabidopsis*, *Brassica*, *Raphanus*, *Sinapis*, *Atropa*, *Capsicum*, *Datura*, *Hyoscyamus*, *Lycopersion*, *Nicotiana*, *Solanum*, *Petunia*, *Digitalis*, *Majorana*, *Cichorium*, *Helianthus*, *Lactuca*, *Bromus*, *Asparagus*, *Antirrhinum*, *Hererocallis*, *Nemesia*, *Pelargonium*,
10 *Panicum*, *Pennisetum*, *Ranunculus*, *Senecio*, *Salpiglossis*, *Cucumis*, *Browaalia*, *Glycine*, *Lolium*, *Zea*, *Triticum*, *Sorghum*, and *Datura*.

Means for regeneration vary from species to species of plants, but generally a suspension of transformed protoplasts containing copies of the heterologous gene is first provided. Callus tissue is formed and shoots may be induced from callus and subsequently rooted. Alternatively, embryo
15 formation can be induced from the protoplast suspension. These embryos germinate as natural embryos to form plants. The culture media will generally contain various amino acids and hormones, such as auxin and cytokinins. It is also advantageous to add glutamic acid and proline to the medium, especially for such species as corn and alfalfa. Shoots and roots normally develop simultaneously. Efficient regeneration will depend on the medium, on the genotype, and on the
20 history of the culture. If these three variables are controlled, then regeneration is fully reproducible and repeatable.

In some plant cell culture systems, the desired protein of the invention may be excreted or alternatively, the protein may be extracted from the whole plant. Where the desired protein of the invention is secreted into the medium, it may be collected. Alternatively, the embryos and
25 embryoless-half seeds or other plant tissue may be mechanically disrupted to release any secreted protein between cells and tissues. The mixture may be suspended in a buffer solution to retrieve soluble proteins. Conventional protein isolation and purification methods will be then used to purify the recombinant protein. Parameters of time, temperature pH, oxygen, and volumes will be adjusted through routine methods to optimize expression and recovery of heterologous protein.

iv. Bacterial Systems

Bacterial expression techniques are known in the art. A bacterial promoter is any DNA sequence capable of binding bacterial RNA polymerase and initiating the downstream (3') transcription of a coding sequence (eg. structural gene) into mRNA. A promoter will have a transcription initiation
5 region which is usually placed proximal to the 5' end of the coding sequence. This transcription initiation region usually includes an RNA polymerase binding site and a transcription initiation site. A bacterial promoter may also have a second domain called an operator, that may overlap an adjacent RNA polymerase binding site at which RNA synthesis begins. The operator permits negative regulated (inducible) transcription, as a gene repressor protein may bind the operator and
10 thereby inhibit transcription of a specific gene. Constitutive expression may occur in the absence of negative regulatory elements, such as the operator. In addition, positive regulation may be achieved by a gene activator protein binding sequence, which, if present is usually proximal (5') to the RNA polymerase binding sequence. An example of a gene activator protein is the catabolite activator protein (CAP), which helps initiate transcription of the lac operon in *Escherichia coli* (*E. coli*) [Raibaud *et al.* (1984) *Annu. Rev. Genet.* 18:173]. Regulated expression may therefore be
15 either positive or negative, thereby either enhancing or reducing transcription.

Sequences encoding metabolic pathway enzymes provide particularly useful promoter sequences. Examples include promoter sequences derived from sugar metabolizing enzymes, such as galactose, lactose (*lac*) [Chang *et al.* (1977) *Nature* 198:1056], and maltose. Additional examples include
20 promoter sequences derived from biosynthetic enzymes such as tryptophan (*trp*) [Goeddel *et al.* (1980) *Nuc. Acids Res.* 8:4057; Yelverton *et al.* (1981) *Nucl. Acids Res.* 9:731; US patent 4,738,921; EP-A-0036776 and EP-A-0121775]. The g-lactamase (*bla*) promoter system [Weissmann (1981) "The cloning of interferon and other mistakes." In *Interferon 3* (ed. I. Gresser)], bacteriophage lambda PL [Shimatake *et al.* (1981) *Nature* 292:128] and T5 [US patent 4,689,406]
25 promoter systems also provide useful promoter sequences.

In addition, synthetic promoters which do not occur in nature also function as bacterial promoters. For example, transcription activation sequences of one bacterial or bacteriophage promoter may be joined with the operon sequences of another bacterial or bacteriophage promoter, creating a synthetic hybrid promoter [US patent 4,551,433]. For example, the *tac* promoter is a hybrid *trp-lac*
30 promoter comprised of both *trp* promoter and *lac* operon sequences that is regulated by the *lac* repressor [Amann *et al.* (1983) *Gene* 25:167; de Boer *et al.* (1983) *Proc. Natl. Acad. Sci.* 80:21].

Furthermore, a bacterial promoter can include naturally occurring promoters of non-bacterial origin that have the ability to bind bacterial RNA polymerase and initiate transcription. A naturally occurring promoter of non-bacterial origin can also be coupled with a compatible RNA polymerase to produce high levels of expression of some genes in prokaryotes. The bacteriophage T7 RNA
5 polymerase/promoter system is an example of a coupled promoter system [Studier *et al.* (1986) *J. Mol. Biol.* 189:113; Tabor *et al.* (1985) *Proc Natl. Acad. Sci.* 82:1074]. In addition, a hybrid promoter can also be comprised of a bacteriophage promoter and an *E. coli* operator region (EPO-A-0 267 851).

In addition to a functioning promoter sequence, an efficient ribosome binding site is also useful for
10 the expression of foreign genes in prokaryotes. In *E. coli*, the ribosome binding site is called the Shine-Dalgarno (SD) sequence and includes an initiation codon (ATG) and a sequence 3-9 nucleotides in length located 3-11 nucleotides upstream of the initiation codon [Shine *et al.* (1975) *Nature* 254:34]. The SD sequence is thought to promote binding of mRNA to the ribosome by the pairing of bases between the SD sequence and the 3' end of *E. coli* 16S rRNA [Steitz *et al.* (1979)
15 "Genetic signals and nucleotide sequences in messenger RNA." In *Biological Regulation and Development: Gene Expression* (ed. R.F. Goldberger)]. To express eukaryotic genes and prokaryotic genes with weak ribosome-binding site [Sambrook *et al.* (1989) "Expression of cloned genes in *Escherichia coli*." In *Molecular Cloning: A Laboratory Manual*].

A DNA molecule may be expressed intracellularly. A promoter sequence may be directly linked
20 with the DNA molecule, in which case the first amino acid at the N-terminus will always be a methionine, which is encoded by the ATG start codon. If desired, methionine at the N-terminus may be cleaved from the protein by *in vitro* incubation with cyanogen bromide or by either *in vivo* or *in vitro* incubation with a bacterial methionine N-terminal peptidase (EPO-A-0 219 237).

Fusion proteins provide an alternative to direct expression. Usually, a DNA sequence encoding the
25 N-terminal portion of an endogenous bacterial protein, or other stable protein, is fused to the 5' end of heterologous coding sequences. Upon expression, this construct will provide a fusion of the two amino acid sequences. For example, the bacteriophage lambda cell gene can be linked at the 5' terminus of a foreign gene and expressed in bacteria. The resulting fusion protein preferably retains a site for a processing enzyme (factor Xa) to cleave the bacteriophage protein from the foreign gene
30 [Nagai *et al.* (1984) *Nature* 309:810]. Fusion proteins can also be made with sequences from the *lacZ* [Jia *et al.* (1987) *Gene* 60:197], *trpE* [Allen *et al.* (1987) *J. Biotechnol.* 5:93; Makoff *et al.*

(1989) *J. Gen. Microbiol.* 135:11], and Chey [EP-A-0 324 647] genes. The DNA sequence at the junction of the two amino acid sequences may or may not encode a cleavable site. Another example is a ubiquitin fusion protein. Such a fusion protein is made with the ubiquitin region that preferably retains a site for a processing enzyme (eg. ubiquitin specific processing-protease) to cleave the ubiquitin from the foreign protein. Through this method, native foreign protein can be isolated [Miller *et al.* (1989) *Bio/Technology* 7:698].

Alternatively, foreign proteins can also be secreted from the cell by creating chimeric DNA molecules that encode a fusion protein comprised of a signal peptide sequence fragment that provides for secretion of the foreign protein in bacteria [US patent 4,336,336]. The signal sequence fragment usually encodes a signal peptide comprised of hydrophobic amino acids which direct the secretion of the protein from the cell. The protein is either secreted into the growth media (gram-positive bacteria) or into the periplasmic space, located between the inner and outer membrane of the cell (gram-negative bacteria). Preferably there are processing sites, which can be cleaved either *in vivo* or *in vitro* encoded between the signal peptide fragment and the foreign gene.

DNA encoding suitable signal sequences can be derived from genes for secreted bacterial proteins, such as the *E. coli* outer membrane protein gene (*ompA*) [Masui *et al.* (1983), in: *Experimental Manipulation of Gene Expression*; Ghrayeb *et al.* (1984) *EMBO J.* 3:2437] and the *E. coli* alkaline phosphatase signal sequence (*phoA*) [Oka *et al.* (1985) *Proc. Natl. Acad. Sci.* 82:7212]. As an additional example, the signal sequence of the alpha-amylase gene from various *Bacillus* strains can be used to secrete heterologous proteins from *B. subtilis* [Palva *et al.* (1982) *Proc. Natl. Acad. Sci. USA* 79:5582; EP-A-0 244 042].

Usually, transcription termination sequences recognized by bacteria are regulatory regions located 3' to the translation stop codon, and thus together with the promoter flank the coding sequence. These sequences direct the transcription of an mRNA which can be translated into the polypeptide encoded by the DNA. Transcription termination sequences frequently include DNA sequences of about 50 nucleotides capable of forming stem loop structures that aid in terminating transcription. Examples include transcription termination sequences derived from genes with strong promoters, such as the *trp* gene in *E. coli* as well as other biosynthetic genes.

Usually, the above described components, comprising a promoter, signal sequence (if desired), coding sequence of interest, and transcription termination sequence, are put together into expression constructs. Expression constructs are often maintained in a replicon, such as an extrachromosomal

element (*eg.* plasmids) capable of stable maintenance in a host, such as bacteria. The replicon will have a replication system, thus allowing it to be maintained in a prokaryotic host either for expression or for cloning and amplification. In addition, a replicon may be either a high or low copy number plasmid. A high copy number plasmid will generally have a copy number ranging from about 5 to about 200, and usually about 10 to about 150. A host containing a high copy number plasmid will preferably contain at least about 10, and more preferably at least about 20 plasmids. Either a high or low copy number vector may be selected, depending upon the effect of the vector and the foreign protein on the host.

Alternatively, the expression constructs can be integrated into the bacterial genome with an integrating vector. Integrating vectors usually contain at least one sequence homologous to the bacterial chromosome that allows the vector to integrate. Integrations appear to result from recombinations between homologous DNA in the vector and the bacterial chromosome. For example, integrating vectors constructed with DNA from various *Bacillus* strains integrate into the *Bacillus* chromosome (EP-A- 0 127 328). Integrating vectors may also be comprised of bacteriophage or transposon sequences.

Usually, extrachromosomal and integrating expression constructs may contain selectable markers to allow for the selection of bacterial strains that have been transformed. Selectable markers can be expressed in the bacterial host and may include genes which render bacteria resistant to drugs such as ampicillin, chloramphenicol, erythromycin, kanamycin (neomycin), and tetracycline [Davies *et al.* (1978) *Annu. Rev. Microbiol.* 32:469]. Selectable markers may also include biosynthetic genes, such as those in the histidine, tryptophan, and leucine biosynthetic pathways.

Alternatively, some of the above described components can be put together in transformation vectors. Transformation vectors are usually comprised of a selectable market that is either maintained in a replicon or developed into an integrating vector, as described above.

Expression and transformation vectors, either extra-chromosomal replicons or integrating vectors, have been developed for transformation into many bacteria. For example, expression vectors have been developed for, *inter alia*, the following bacteria: *Bacillus subtilis* [Palva *et al.* (1982) *Proc. Natl. Acad. Sci. USA* 79:5582; EP-A-0 036 259 and EP-A-0 063 953; WO 84/04541], *Escherichia coli* [Shimatake *et al.* (1981) *Nature* 292:128; Amann *et al.* (1985) *Gene* 40:183; Studier *et al.* (1986) *J. Mol. Biol.* 189:113; EP-A-0 036 776, EP-A-0 136 829 and EP-A-0 136 907],

Streptococcus cremoris [Powell *et al.* (1988) *Appl. Environ. Microbiol.* 54:655]; *Streptococcus lividans* [Powell *et al.* (1988) *Appl. Environ. Microbiol.* 54:655], *Streptomyces lividans* [US patent 4,745,056].

Methods of introducing exogenous DNA into bacterial hosts are well-known in the art, and usually include either the transformation of bacteria treated with CaCl_2 or other agents, such as divalent cations and DMSO. DNA can also be introduced into bacterial cells by electroporation. Transformation procedures usually vary with the bacterial species to be transformed. See *eg.* [Masson *et al.* (1989) *FEMS Microbiol. Lett.* 60:273; Palva *et al.* (1982) *Proc. Natl. Acad. Sci. USA* 79:5582; EP-A-0 036 259 and EP-A-0 063 953; WO 84/04541, *Bacillus*], [Miller *et al.* (1988) *Proc. Natl. Acad. Sci.* 85:856; Wang *et al.* (1990) *J. Bacteriol.* 172:949, *Campylobacter*], [Cohen *et al.* (1973) *Proc. Natl. Acad. Sci.* 69:2110; Dower *et al.* (1988) *Nucleic Acids Res.* 16:6127; Kushner (1978) "An improved method for transformation of *Escherichia coli* with ColE1-derived plasmids. In *Genetic Engineering: Proceedings of the International Symposium on Genetic Engineering* (eds. H.W. Boyer and S. Nicosia); Mandel *et al.* (1970) *J. Mol. Biol.* 53:159; Taketo (1988) *Biochim. Biophys. Acta* 949:318; *Escherichia*], [Chassy *et al.* (1987) *FEMS Microbiol. Lett.* 44:173 *Lactobacillus*]; [Fiedler *et al.* (1988) *Anal. Biochem* 170:38, *Pseudomonas*]; [Augustin *et al.* (1990) *FEMS Microbiol. Lett.* 66:203, *Staphylococcus*], [Barany *et al.* (1980) *J. Bacteriol.* 144:698; Harlander (1987) "Transformation of *Streptococcus lactis* by electroporation, in: *Streptococcal Genetics* (ed. J. Ferretti and R. Curtiss III); Perry *et al.* (1981) *Infect. Immun.* 32:1295; Powell *et al.* (1988) *Appl. Environ. Microbiol.* 54:655; Somkuti *et al.* (1987) *Proc. 4th Evr. Cong. Biotechnology* 1:412, *Streptococcus*].

v. Yeast Expression

Yeast expression systems are also known to one of ordinary skill in the art. A yeast promoter is any DNA sequence capable of binding yeast RNA polymerase and initiating the downstream (3') transcription of a coding sequence (*eg.* structural gene) into mRNA. A promoter will have a transcription initiation region which is usually placed proximal to the 5' end of the coding sequence. This transcription initiation region usually includes an RNA polymerase binding site (the "TATA Box") and a transcription initiation site. A yeast promoter may also have a second domain called an upstream activator sequence (UAS), which, if present, is usually distal to the structural gene. The UAS permits regulated (inducible) expression. Constitutive expression occurs in the absence

of a UAS. Regulated expression may be either positive or negative, thereby either enhancing or reducing transcription.

Yeast is a fermenting organism with an active metabolic pathway, therefore sequences encoding enzymes in the metabolic pathway provide particularly useful promoter sequences. Examples
5 include alcohol dehydrogenase (ADH) (EP-A-0 284 044), enolase, glucokinase, glucose-6-phosphate isomerase, glyceraldehyde-3-phosphate-dehydrogenase (GAP or GAPDH), hexokinase, phosphofructokinase, 3-phosphoglycerate mutase, and pyruvate kinase (PyK) (EPO-A-0 329 203). The yeast *PHO5* gene, encoding acid phosphatase, also provides useful promoter sequences [Myanohara *et al.* (1983) *Proc. Natl. Acad. Sci. USA* 80:1].

10 In addition, synthetic promoters which do not occur in nature also function as yeast promoters. For example, UAS sequences of one yeast promoter may be joined with the transcription activation region of another yeast promoter, creating a synthetic hybrid promoter. Examples of such hybrid promoters include the ADH regulatory sequence linked to the GAP transcription activation region (US Patent Nos. 4,876,197 and 4,880,734). Other examples of hybrid promoters include promoters
15 which consist of the regulatory sequences of either the *ADH2*, *GAL4*, *GAL10*, OR *PHO5* genes, combined with the transcriptional activation region of a glycolytic enzyme gene such as GAP or PyK (EP-A-0 164 556). Furthermore, a yeast promoter can include naturally occurring promoters of non-yeast origin that have the ability to bind yeast RNA polymerase and initiate transcription. Examples of such promoters include, *inter alia*, [Cohen *et al.* (1980) *Proc. Natl. Acad. Sci. USA*
20 77:1078; Henikoff *et al.* (1981) *Nature* 283:835; Hollenberg *et al.* (1981) *Curr. Topics Microbiol. Immunol.* 96:119; Hollenberg *et al.* (1979) "The Expression of Bacterial Antibiotic Resistance Genes in the Yeast *Saccharomyces cerevisiae*," in: *Plasmids of Medical, Environmental and Commercial Importance* (eds. K.N. Timmis and A. Puhler); Mercerau-Puigalon *et al.* (1980) *Gene* 11:163; Panthier *et al.* (1980) *Curr. Genet.* 2:109;].

25 A DNA molecule may be expressed intracellularly in yeast. A promoter sequence may be directly linked with the DNA molecule, in which case the first amino acid at the N-terminus of the recombinant protein will always be a methionine, which is encoded by the ATG start codon. If desired, methionine at the N-terminus may be cleaved from the protein by *in vitro* incubation with cyanogen bromide.

Fusion proteins provide an alternative for yeast expression systems, as well as in mammalian, baculovirus, and bacterial expression systems. Usually, a DNA sequence encoding the N-terminal portion of an endogenous yeast protein, or other stable protein, is fused to the 5' end of heterologous coding sequences. Upon expression, this construct will provide a fusion of the two amino acid sequences. For example, the yeast or human superoxide dismutase (SOD) gene, can be linked at the 5' terminus of a foreign gene and expressed in yeast. The DNA sequence at the junction of the two amino acid sequences may or may not encode a cleavable site. See *eg.* EP-A-0 196 056. Another example is a ubiquitin fusion protein. Such a fusion protein is made with the ubiquitin region that preferably retains a site for a processing enzyme (*eg.* ubiquitin-specific processing protease) to cleave the ubiquitin from the foreign protein. Through this method, therefore, native foreign protein can be isolated (*eg.* WO88/024066).

Alternatively, foreign proteins can also be secreted from the cell into the growth media by creating chimeric DNA molecules that encode a fusion protein comprised of a leader sequence fragment that provide for secretion in yeast of the foreign protein. Preferably, there are processing sites encoded between the leader fragment and the foreign gene that can be cleaved either *in vivo* or *in vitro*. The leader sequence fragment usually encodes a signal peptide comprised of hydrophobic amino acids which direct the secretion of the protein from the cell.

DNA encoding suitable signal sequences can be derived from genes for secreted yeast proteins, such as the yeast invertase gene (EP-A-0 012 873; JPO. 62,096,086) and the A-factor gene (US patent 4,588,684). Alternatively, leaders of non-yeast origin, such as an interferon leader, exist that also provide for secretion in yeast (EP-A-0 060 057).

A preferred class of secretion leaders are those that employ a fragment of the yeast alpha-factor gene, which contains both a "pre" signal sequence, and a "pro" region. The types of alpha-factor fragments that can be employed include the full-length pre-pro alpha factor leader (about 83 amino acid residues) as well as truncated alpha-factor leaders (usually about 25 to about 50 amino acid residues) (US Patents 4,546,083 and 4,870,008; EP-A-0 324 274). Additional leaders employing an alpha-factor leader fragment that provides for secretion include hybrid alpha-factor leaders made with a presequence of a first yeast, but a pro-region from a second yeast alphafactor. (*eg.* see WO 89/02463.)

Usually, transcription termination sequences recognized by yeast are regulatory regions located 3' to the translation stop codon, and thus together with the promoter flank the coding sequence. These sequences direct the transcription of an mRNA which can be translated into the polypeptide encoded by the DNA. Examples of transcription terminator sequence and other yeast-recognized termination sequences, such as those coding for glycolytic enzymes.

Usually, the above described components, comprising a promoter, leader (if desired), coding sequence of interest, and transcription termination sequence, are put together into expression constructs. Expression constructs are often maintained in a replicon, such as an extrachromosomal element (eg. plasmids) capable of stable maintenance in a host, such as yeast or bacteria. The replicon may have two replication systems, thus allowing it to be maintained, for example, in yeast for expression and in a prokaryotic host for cloning and amplification. Examples of such yeast-bacteria shuttle vectors include YEp24 [Botstein *et al.* (1979) *Gene* 8:17-24], pCI/1 [Brake *et al.* (1984) *Proc. Natl. Acad. Sci USA* 81:4642-4646], and YRp17 [Stinchcomb *et al.* (1982) *J. Mol. Biol.* 158:157]. In addition, a replicon may be either a high or low copy number plasmid. A high copy number plasmid will generally have a copy number ranging from about 5 to about 200, and usually about 10 to about 150. A host containing a high copy number plasmid will preferably have at least about 10, and more preferably at least about 20. Enter a high or low copy number vector may be selected, depending upon the effect of the vector and the foreign protein on the host. See eg. Brake *et al.*, *supra*.

Alternatively, the expression constructs can be integrated into the yeast genome with an integrating vector. Integrating vectors usually contain at least one sequence homologous to a yeast chromosome that allows the vector to integrate, and preferably contain two homologous sequences flanking the expression construct. Integrations appear to result from recombinations between homologous DNA in the vector and the yeast chromosome [Orr-Weaver *et al.* (1983) *Methods in Enzymol.* 101:228-245]. An integrating vector may be directed to a specific locus in yeast by selecting the appropriate homologous sequence for inclusion in the vector. See Orr-Weaver *et al.*, *supra*. One or more expression construct may integrate, possibly affecting levels of recombinant protein produced [Rine *et al.* (1983) *Proc. Natl. Acad. Sci. USA* 80:6750]. The chromosomal sequences included in the vector can occur either as a single segment in the vector, which results in the integration of the entire vector, or two segments homologous to adjacent segments in the

chromosome and flanking the expression construct in the vector, which can result in the stable integration of only the expression construct.

Usually, extrachromosomal and integrating expression constructs may contain selectable markers to allow for the selection of yeast strains that have been transformed. Selectable markers may include biosynthetic genes that can be expressed in the yeast host, such as *ADE2*, *HIS4*, *LEU2*, *TRP1*, and *ALG7*, and the G418 resistance gene, which confer resistance in yeast cells to tunicamycin and G418, respectively. In addition, a suitable selectable marker may also provide yeast with the ability to grow in the presence of toxic compounds, such as metal. For example, the presence of *CUP1* allows yeast to grow in the presence of copper ions [Butt *et al.* (1987) *Microbiol. Rev.* 51:351].

Alternatively, some of the above described components can be put together into transformation vectors. Transformation vectors are usually comprised of a selectable marker that is either maintained in a replicon or developed into an integrating vector, as described above.

Expression and transformation vectors, either extrachromosomal replicons or integrating vectors, have been developed for transformation into many yeasts. For example, expression vectors have been developed for, *inter alia*, the following yeasts: *Candida albicans* [Kurtz, *et al.* (1986) *Mol. Cell. Biol.* 6:142], *Candida maltosa* [Kunze, *et al.* (1985) *J. Basic Microbiol.* 25:141], *Hansenula polymorpha* [Gleeson, *et al.* (1986) *J. Gen. Microbiol.* 132:3459; Roggenkamp *et al.* (1986) *Mol. Gen. Genet.* 202:302], *Kluyveromyces fragilis* [Das, *et al.* (1984) *J. Bacteriol.* 158:1165], *Kluyveromyces lactis* [De Louvencourt *et al.* (1983) *J. Bacteriol.* 154:737; Van den Berg *et al.* (1990) *Bio/Technology* 8:135], *Pichia guilliermondii* [Kunze *et al.* (1985) *J. Basic Microbiol.* 25:141], *Pichia pastoris* [Cregg, *et al.* (1985) *Mol. Cell. Biol.* 5:3376; US Patent Nos. 4,837,148 and 4,929,555], *Saccharomyces cerevisiae* [Hinnen *et al.* (1978) *Proc. Natl. Acad. Sci. USA* 75:1929; Ito *et al.* (1983) *J. Bacteriol.* 153:163], *Schizosaccharomyces pombe* [Beach and Nurse (1981) *Nature* 300:706], and *Yarrowia lipolytica* [Davidow, *et al.* (1985) *Curr. Genet.* 10:380471 Gaillardin, *et al.* (1985) *Curr. Genet.* 10:49].

Methods of introducing exogenous DNA into yeast hosts are well-known in the art, and usually include either the transformation of spheroplasts or of intact yeast cells treated with alkali cations. Transformation procedures usually vary with the yeast species to be transformed. See *eg.* [Kurtz *et al.* (1986) *Mol. Cell. Biol.* 6:142; Kunze *et al.* (1985) *J. Basic Microbiol.* 25:141; *Candida*];

[Gleeson *et al.* (1986) *J. Gen. Microbiol.* 132:3459; Roggenkamp *et al.* (1986) *Mol. Gen. Genet.* 202:302; Hansenula]; [Das *et al.* (1984) *J. Bacteriol.* 158:1165; De Louvencourt *et al.* (1983) *J. Bacteriol.* 154:1165; Van den Berg *et al.* (1990) *Bio/Technology* 8:135; Kluyveromyces]; [Cregg *et al.* (1985) *Mol. Cell. Biol.* 5:3376; Kunze *et al.* (1985) *J. Basic Microbiol.* 25:141; US Patent
5 Nos. 4,837,148 and 4,929,555; Pichia]; [Hinnen *et al.* (1978) *Proc. Natl. Acad. Sci. USA* 75:1929; Ito *et al.* (1983) *J. Bacteriol.* 153:163 Saccharomyces]; [Beach and Nurse (1981) *Nature* 300:706; Schizosaccharomyces]; [Davidow *et al.* (1985) *Curr. Genet.* 10:39; Gaillardin *et al.* (1985) *Curr. Genet.* 10:49; Yarrowia].

Antibodies

10 As used herein, the term “antibody” refers to a polypeptide or group of polypeptides composed of at least one antibody combining site. An “antibody combining site” is the three-dimensional binding space with an internal surface shape and charge distribution complementary to the features of an epitope of an antigen, which allows a binding of the antibody with the antigen. “Antibody” includes, for example, vertebrate antibodies, hybrid antibodies, chimeric antibodies, humanised
15 antibodies, altered antibodies, univalent antibodies, Fab proteins, and single domain antibodies.

Antibodies against the proteins of the invention are useful for affinity chromatography, immunoassays, and distinguishing/identifying Neisserial proteins.

Antibodies to the proteins of the invention, both polyclonal and monoclonal, may be prepared by conventional methods. In general, the protein is first used to immunize a suitable animal, preferably
20 a mouse, rat, rabbit or goat. Rabbits and goats are preferred for the preparation of polyclonal sera due to the volume of serum obtainable, and the availability of labeled anti-rabbit and anti-goat antibodies. Immunization is generally performed by mixing or emulsifying the protein in saline, preferably in an adjuvant such as Freund’s complete adjuvant, and injecting the mixture or emulsion parenterally (generally subcutaneously or intramuscularly). A dose of 50-200 µg/injection
25 is typically sufficient. Immunization is generally boosted 2-6 weeks later with one or more injections of the protein in saline, preferably using Freund’s incomplete adjuvant. One may alternatively generate antibodies by *in vitro* immunization using methods known in the art, which for the purposes of this invention is considered equivalent to *in vivo* immunization. Polyclonal antisera is obtained by bleeding the immunized animal into a glass or plastic container, incubating
30 the blood at 25°C for one hour, followed by incubating at 4°C for 2-18 hours. The serum is

recovered by centrifugation (eg. 1,000g for 10 minutes). About 20-50 ml per bleed may be obtained from rabbits.

Monoclonal antibodies are prepared using the standard method of Kohler & Milstein [*Nature* (1975) 256:495-96], or a modification thereof. Typically, a mouse or rat is immunized as described
5 above. However, rather than bleeding the animal to extract serum, the spleen (and optionally several large lymph nodes) is removed and dissociated into single cells. If desired, the spleen cells may be screened (after removal of nonspecifically adherent cells) by applying a cell suspension to a plate or well coated with the protein antigen. B-cells expressing membrane-bound immunoglobulin specific for the antigen bind to the plate, and are not rinsed away with the rest of
10 the suspension. Resulting B-cells, or all dissociated spleen cells, are then induced to fuse with myeloma cells to form hybridomas, and are cultured in a selective medium (eg. hypoxanthine, aminopterin, thymidine medium, "HAT"). The resulting hybridomas are plated by limiting dilution, and are assayed for the production of antibodies which bind specifically to the immunizing antigen (and which do not bind to unrelated antigens). The selected MAb-secreting hybridomas are then
15 cultured either *in vitro* (eg. in tissue culture bottles or hollow fiber reactors), or *in vivo* (as ascites in mice).

If desired, the antibodies (whether polyclonal or monoclonal) may be labeled using conventional techniques. Suitable labels include fluorophores, chromophores, radioactive atoms (particularly ³²P and ¹²⁵I), electron-dense reagents, enzymes, and ligands having specific binding partners. Enzymes
20 are typically detected by their activity. For example, horseradish peroxidase is usually detected by its ability to convert 3,3',5,5'-tetramethylbenzidine (TMB) to a blue pigment, quantifiable with a spectrophotometer. "Specific binding partner" refers to a protein capable of binding a ligand molecule with high specificity, as for example in the case of an antigen and a monoclonal antibody specific therefor. Other specific binding partners include biotin and avidin or streptavidin, IgG and protein A,
25 and the numerous receptor-ligand couples known in the art. It should be understood that the above description is not meant to categorize the various labels into distinct classes, as the same label may serve in several different modes. For example, ¹²⁵I may serve as a radioactive label or as an electron-dense reagent. HRP may serve as enzyme or as antigen for a MAb. Further, one may combine various labels for desired effect. For example, MAbs and avidin also require labels in the practice of
30 this invention: thus, one might label a MAb with biotin, and detect its presence with avidin labeled with ¹²⁵I, or with an anti-biotin MAb labeled with HRP. Other permutations and possibilities will be

readily apparent to those of ordinary skill in the art, and are considered as equivalents within the scope of the instant invention.

Pharmaceutical Compositions

Pharmaceutical compositions can comprise either polypeptides, antibodies, or nucleic acid of the invention. The pharmaceutical compositions will comprise a therapeutically effective amount of
5 either polypeptides, antibodies, or polynucleotides of the claimed invention.

The term "therapeutically effective amount" as used herein refers to an amount of a therapeutic agent to treat, ameliorate, or prevent a desired disease or condition, or to exhibit a detectable therapeutic or preventative effect. The effect can be detected by, for example, chemical markers or
10 antigen levels. Therapeutic effects also include reduction in physical symptoms, such as decreased body temperature. The precise effective amount for a subject will depend upon the subject's size and health, the nature and extent of the condition, and the therapeutics or combination of therapeutics selected for administration. Thus, it is not useful to specify an exact effective amount in advance. However, the effective amount for a given situation can be determined by routine
15 experimentation and is within the judgement of the clinician.

For purposes of the present invention, an effective dose will be from about 0.01 mg/ kg to 50 mg/kg or 0.05 mg/kg to about 10 mg/kg of the DNA constructs in the individual to which it is administered.

A pharmaceutical composition can also contain a pharmaceutically acceptable carrier. The term "pharmaceutically acceptable carrier" refers to a carrier for administration of a therapeutic agent, such
20 as antibodies or a polypeptide, genes, and other therapeutic agents. The term refers to any pharmaceutical carrier that does not itself induce the production of antibodies harmful to the individual receiving the composition, and which may be administered without undue toxicity. Suitable carriers may be large, slowly metabolized macromolecules such as proteins, polysaccharides, polylactic acids, polyglycolic acids, polymeric amino acids, amino acid copolymers, and inactive virus
25 particles. Such carriers are well known to those of ordinary skill in the art.

Pharmaceutically acceptable salts can be used therein, for example, mineral acid salts such as hydrochlorides, hydrobromides, phosphates, sulfates, and the like; and the salts of organic acids such as acetates, propionates, malonates, benzoates, and the like. A thorough discussion of pharmaceutically acceptable excipients is available in Remington's Pharmaceutical Sciences (Mack
30 Pub. Co., N.J. 1991).

Pharmaceutically acceptable carriers in therapeutic compositions may contain liquids such as water, saline, glycerol and ethanol. Additionally, auxiliary substances, such as wetting or emulsifying agents, pH buffering substances, and the like, may be present in such vehicles. Typically, the therapeutic compositions are prepared as injectables, either as liquid solutions or suspensions; solid forms suitable for solution in, or suspension in, liquid vehicles prior to injection may also be prepared. Liposomes are included within the definition of a pharmaceutically acceptable carrier.

Delivery Methods

Once formulated, the compositions of the invention can be administered directly to the subject. The subjects to be treated can be animals; in particular, human subjects can be treated.

Direct delivery of the compositions will generally be accomplished by injection, either subcutaneously, intraperitoneally, intravenously or intramuscularly or delivered to the interstitial space of a tissue. The compositions can also be administered into a lesion. Other modes of administration include oral and pulmonary administration, suppositories, and transdermal or transcutaneous applications (*eg.* see WO98/20734), needles, and gene guns or hyposprays. Dosage treatment may be a single dose schedule or a multiple dose schedule.

Vaccines

Vaccines according to the invention may either be prophylactic (*ie.* to prevent infection) or therapeutic (*ie.* to treat disease after infection).

Such vaccines comprise immunising antigen(s), immunogen(s), polypeptide(s), protein(s) or nucleic acid, usually in combination with "pharmaceutically acceptable carriers," which include any carrier that does not itself induce the production of antibodies harmful to the individual receiving the composition. Suitable carriers are typically large, slowly metabolized macromolecules such as proteins, polysaccharides, polylactic acids, polyglycolic acids, polymeric amino acids, amino acid copolymers, lipid aggregates (such as oil droplets or liposomes), and inactive virus particles. Such carriers are well known to those of ordinary skill in the art. Additionally, these carriers may function as immunostimulating agents ("adjuvants"). Furthermore, the antigen or immunogen may be conjugated to a bacterial toxoid, such as a toxoid from diphtheria, tetanus, cholera, *H. pylori*, *etc.* pathogens.

Preferred adjuvants to enhance effectiveness of the composition include, but are not limited to: (1) aluminum salts (alum), such as aluminum hydroxide, aluminum phosphate, aluminum sulfate, *etc.*; (2) oil-in-water emulsion formulations (with or without other specific immunostimulating agents

such as muramyl peptides (see below) or bacterial cell wall components), such as for example (a) MF59™ (WO 90/14837; Chapter 10 in *Vaccine design: the subunit and adjuvant approach*, eds. Powell & Newman, Plenum Press 1995), containing 5% Squalene, 0.5% Tween 80, and 0.5% Span 85 (optionally containing various amounts of MTP-PE (see below), although not required) formulated into submicron particles using a microfluidizer such as Model 110Y microfluidizer (Microfluidics, Newton, MA), (b) SAF, containing 10% Squalane, 0.4% Tween 80, 5% pluronic-blocked polymer L121, and thr-MDP (see below) either microfluidized into a submicron emulsion or vortexed to generate a larger particle size emulsion, and (c) Ribi™ adjuvant system (RAS), (Ribi Immunochem, Hamilton, MT) containing 2% Squalene, 0.2% Tween 80, and one or more bacterial cell wall components from the group consisting of monophosphorylipid A (MPL), trehalose dimycolate (TDM), and cell wall skeleton (CWS), preferably MPL + CWS (Detox™); (3) saponin adjuvants, such as Stimulon™ (Cambridge Bioscience, Worcester, MA) may be used or particles generated therefrom such as ISCOMs (immunostimulating complexes); (4) Complete Freund's Adjuvant (CFA) and Incomplete Freund's Adjuvant (IFA); (5) cytokines, such as interleukins (*eg.* IL-1, IL-2, IL-4, IL-5, IL-6, IL-7, IL-12, *etc.*), interferons (*eg.* gamma interferon), macrophage colony stimulating factor (M-CSF), tumor necrosis factor (TNF), *etc.*; and (6) other substances that act as immunostimulating agents to enhance the effectiveness of the composition. Alum and MF59™ are preferred.

As mentioned above, muramyl peptides include, but are not limited to, N-acetyl-muramyl-L-threonyl-D-isoglutamine (thr-MDP), N-acetyl-normuramyl-L-alanyl-D-isoglutamine (nor-MDP), N-acetylmuramyl-L-alanyl-D-isoglutaminyl-L-alanine-2-(1'-2'-dipalmitoyl-*sn*-glycero-3-hydroxyphosphoryloxy)-ethylamine (MTP-PE), *etc.*

The immunogenic compositions (*eg.* the immunising antigen/immunogen/polypeptide/protein/nucleic acid, pharmaceutically acceptable carrier, and adjuvant) typically will contain diluents, such as water, saline, glycerol, ethanol, *etc.* Additionally, auxiliary substances, such as wetting or emulsifying agents, pH buffering substances, and the like, may be present in such vehicles.

Typically, the immunogenic compositions are prepared as injectables, either as liquid solutions or suspensions; solid forms suitable for solution in, or suspension in, liquid vehicles prior to injection may also be prepared. The preparation also may be emulsified or encapsulated in liposomes for enhanced adjuvant effect, as discussed above under pharmaceutically acceptable carriers.

Immunogenic compositions used as vaccines comprise an immunologically effective amount of the antigenic or immunogenic polypeptides, as well as any other of the above-mentioned components, as needed. By "immunologically effective amount", it is meant that the administration of that amount to an individual, either in a single dose or as part of a series, is effective for treatment or
5 prevention. This amount varies depending upon the health and physical condition of the individual to be treated, the taxonomic group of individual to be treated (*eg.* nonhuman primate, primate, *etc.*), the capacity of the individual's immune system to synthesize antibodies, the degree of protection desired, the formulation of the vaccine, the treating doctor's assessment of the medical situation, and other relevant factors. It is expected that the amount will fall in a relatively broad range that
10 can be determined through routine trials.

The immunogenic compositions are conventionally administered parenterally, *eg.* by injection, either subcutaneously, intramuscularly, or transdermally/transcutaneously (*eg.* WO98/20734). Additional formulations suitable for other modes of administration include oral and pulmonary formulations, suppositories, and transdermal applications. Dosage treatment may be a single dose
15 schedule or a multiple dose schedule. The vaccine may be administered in conjunction with other immunoregulatory agents.

As an alternative to protein-based vaccines, DNA vaccination may be employed [*eg.* Robinson & Torres (1997) *Seminars in Immunology* 9:271-283; Donnelly *et al.* (1997) *Annu Rev Immunol* 15:617-648; see later herein].

20 Gene Delivery Vehicles

Gene therapy vehicles for delivery of constructs including a coding sequence of a therapeutic of the invention, to be delivered to the mammal for expression in the mammal, can be administered either locally or systemically. These constructs can utilize viral or non-viral vector approaches in *in vivo* or *ex vivo* modality. Expression of such coding sequence can be induced using endogenous
25 mammalian or heterologous promoters. Expression of the coding sequence *in vivo* can be either constitutive or regulated.

The invention includes gene delivery vehicles capable of expressing the contemplated nucleic acid sequences. The gene delivery vehicle is preferably a viral vector and, more preferably, a retroviral, adenoviral, adeno-associated viral (AAV), herpes viral, or alphavirus vector. The viral vector can
30 also be an astrovirus, coronavirus, orthomyxovirus, papovavirus, paramyxovirus, parvovirus,

picornavirus, poxvirus, or togavirus viral vector. See generally, Jolly (1994) *Cancer Gene Therapy* 1:51-64; Kimura (1994) *Human Gene Therapy* 5:845-852; Connelly (1995) *Human Gene Therapy* 6:185-193; and Kaplitt (1994) *Nature Genetics* 6:148-153.

5 Retroviral vectors are well known in the art and we contemplate that any retroviral gene therapy vector is employable in the invention, including B, C and D type retroviruses, xenotropic retroviruses (for example, NZB-X1, NZB-X2 and NZB9-1 (see O'Neill (1985) *J. Virol.* 53:160) polytropic retroviruses eg. MCF and MCF-MLV (see Kelly (1983) *J. Virol.* 45:291), spumaviruses and lentiviruses. See RNA Tumor Viruses, Second Edition, Cold Spring Harbor Laboratory, 1985.

10 Portions of the retroviral gene therapy vector may be derived from different retroviruses. For example, retrovector LTRs may be derived from a Murine Sarcoma Virus, a tRNA binding site from a Rous Sarcoma Virus, a packaging signal from a Murine Leukemia Virus, and an origin of second strand synthesis from an Avian Leukosis Virus.

15 These recombinant retroviral vectors may be used to generate transduction competent retroviral vector particles by introducing them into appropriate packaging cell lines (see US patent 5,591,624). Retrovirus vectors can be constructed for site-specific integration into host cell DNA by incorporation of a chimeric integrase enzyme into the retroviral particle (see WO96/37626). It is preferable that the recombinant viral vector is a replication defective recombinant virus.

20 Packaging cell lines suitable for use with the above-described retrovirus vectors are well known in the art, are readily prepared (see WO95/30763 and WO92/05266), and can be used to create producer cell lines (also termed vector cell lines or "VCLs") for the production of recombinant vector particles. Preferably, the packaging cell lines are made from human parent cells (eg. HT1080 cells) or mink parent cell lines, which eliminates inactivation in human serum.

25 Preferred retroviruses for the construction of retroviral gene therapy vectors include Avian Leukosis Virus, Bovine Leukemia, Virus, Murine Leukemia Virus, Mink-Cell Focus-Inducing Virus, Murine Sarcoma Virus, Reticuloendotheliosis Virus and Rous Sarcoma Virus. Particularly preferred Murine Leukemia Viruses include 4070A and 1504A (Hartley and Rowe (1976) *J Virol* 19:19-25), Abelson (ATCC No. VR-999), Friend (ATCC No. VR-245), Graffi, Gross (ATCC No. VR-590), Kirsten, Harvey Sarcoma Virus and Rauscher (ATCC No. VR-998) and Moloney Murine Leukemia Virus (ATCC No. VR-190). Such retroviruses may be obtained from depositories or

collections such as the American Type Culture Collection ("ATCC") in Rockville, Maryland or isolated from known sources using commonly available techniques.

Exemplary known retroviral gene therapy vectors employable in this invention include those described in patent applications GB2200651, EP0415731, EP0345242, EP0334301, WO89/02468; 5 WO89/05349, WO89/09271, WO90/02806, WO90/07936, WO94/03622, WO93/25698, WO93/25234, WO93/11230, WO93/10218, WO91/02805, WO91/02825, WO95/07994, US 5,219,740, US 4,405,712, US 4,861,719, US 4,980,289, US 4,777,127, US 5,591,624. See also Vile (1993) *Cancer Res* 53:3860-3864; Vile (1993) *Cancer Res* 53:962-967; Ram (1993) *Cancer Res* 53 (1993) 83-88; Takamiya (1992) *J Neurosci Res* 33:493-503; Baba (1993) *J Neurosurg* 10 79:729-735; Mann (1983) *Cell* 33:153; Cane (1984) *Proc Natl Acad Sci* 81:6349; and Miller (1990) *Human Gene Therapy* 1.

Human adenoviral gene therapy vectors are also known in the art and employable in this invention. See, for example, Berkner (1988) *Biotechniques* 6:616 and Rosenfeld (1991) *Science* 252:431, and WO93/07283, WO93/06223, and WO93/07282. Exemplary known adenoviral gene therapy vectors 15 employable in this invention include those described in the above referenced documents and in WO94/12649, WO93/03769, WO93/19191, WO94/28938, WO95/11984, WO95/00655, WO95/27071, WO95/29993, WO95/34671, WO96/05320, WO94/08026, WO94/11506, WO93/06223, WO94/24299, WO95/14102, WO95/24297, WO95/02697, WO94/28152, WO94/24299, WO95/09241, WO95/25807, WO95/05835, WO94/18922 and WO95/09654.

20 Alternatively, administration of DNA linked to killed adenovirus as described in Curiel (1992) *Hum. Gene Ther.* 3:147-154 may be employed. The gene delivery vehicles of the invention also include adenovirus associated virus (AAV) vectors. Leading and preferred examples of such vectors for use in this invention are the AAV-2 based vectors disclosed in Srivastava, WO93/09239. Most preferred AAV vectors comprise the two AAV inverted terminal repeats in 25 which the native D-sequences are modified by substitution of nucleotides, such that at least 5 native nucleotides and up to 18 native nucleotides, preferably at least 10 native nucleotides up to 18 native nucleotides, most preferably 10 native nucleotides are retained and the remaining nucleotides of the D-sequence are deleted or replaced with non-native nucleotides. The native D-sequences of the AAV inverted terminal repeats are sequences of 20 consecutive nucleotides in each AAV inverted 30 terminal repeat (*ie.* there is one sequence at each end) which are not involved in HP formation. The non-native replacement nucleotide may be any nucleotide other than the nucleotide found in the

native D-sequence in the same position. Other employable exemplary AAV vectors are pWP-19, pWN-1, both of which are disclosed in Nahreini (1993) *Gene* 124:257-262. Another example of such an AAV vector is psub201 (see Samulski (1987) *J. Virol.* 61:3096). Another exemplary AAV vector is the Double-D ITR vector. Construction of the Double-D ITR vector is disclosed in US Patent 5,478,745. Still other vectors are those disclosed in Carter US Patent 4,797,368 and Muzyczka US Patent 5,139,941, Chartejee US Patent 5,474,935, and Kotin WO94/288157. Yet a further example of an AAV vector employable in this invention is SSV9AFABTKneo, which contains the AFP enhancer and albumin promoter and directs expression predominantly in the liver. Its structure and construction are disclosed in Su (1996) *Human Gene Therapy* 7:463-470.

5 Additional AAV gene therapy vectors are described in US 5,354,678, US 5,173,414, US 5,139,941, and US 5,252,479.

The gene therapy vectors of the invention also include herpes vectors. Leading and preferred examples are herpes simplex virus vectors containing a sequence encoding a thymidine kinase polypeptide such as those disclosed in US 5,288,641 and EP0176170 (Roizman). Additional exemplary herpes simplex virus vectors include HFEM/ICP6-LacZ disclosed in WO95/04139 (Wistar Institute), pHSVlac described in Geller (1988) *Science* 241:1667-1669 and in WO90/09441 and WO92/07945, HSV Us3::pgC-lacZ described in Fink (1992) *Human Gene Therapy* 3:11-19 and HSV 7134, 2 RH 105 and GAL4 described in EP 0453242 (Breakefield), and those deposited with the ATCC as accession numbers ATCC VR-977 and ATCC VR-260.

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Also contemplated are alpha virus gene therapy vectors that can be employed in this invention. Preferred alpha virus vectors are Sindbis viruses vectors. Togaviruses, Semliki Forest virus (ATCC VR-67; ATCC VR-1247), Middleberg virus (ATCC VR-370), Ross River virus (ATCC VR-373; ATCC VR-1246), Venezuelan equine encephalitis virus (ATCC VR923; ATCC VR-1250; ATCC VR-1249; ATCC VR-532), and those described in US patents 5,091,309, 5,217,879, and WO92/10578. More particularly, those alpha virus vectors described in US Serial No. 08/405,627, filed March 15, 1995, WO94/21792, WO92/10578, WO95/07994, US 5,091,309 and US 5,217,879 are employable. Such alpha viruses may be obtained from depositories or collections such as the ATCC in Rockville, Maryland or isolated from known sources using commonly available techniques. Preferably, alphavirus vectors with reduced cytotoxicity are used (see USSN 08/679640).

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DNA vector systems such as eukaryotic layered expression systems are also useful for expressing the nucleic acids of the invention. See WO95/07994 for a detailed description of eukaryotic layered expression systems. Preferably, the eukaryotic layered expression systems of the invention are derived from alphavirus vectors and most preferably from Sindbis viral vectors.

- 5 Other viral vectors suitable for use in the present invention include those derived from poliovirus, for example ATCC VR-58 and those described in Evans, *Nature* 339 (1989) 385 and Sabin (1973) *J. Biol. Standardization* 1:115; rhinovirus, for example ATCC VR-1110 and those described in Arnold (1990) *J Cell Biochem* L401; pox viruses such as canary pox virus or vaccinia virus, for example ATCC VR-111 and ATCC VR-2010 and those described in Fisher-Hoch (1989) *Proc Natl Acad Sci* 86:317;
- 10 Flexner (1989) *Ann NY Acad Sci* 569:86, Flexner (1990) *Vaccine* 8:17; in US 4,603,112 and US 4,769,330 and WO89/01973; SV40 virus, for example ATCC VR-305 and those described in Mulligan (1979) *Nature* 277:108 and Madzak (1992) *J Gen Virol* 73:1533; influenza virus, for example ATCC VR-797 and recombinant influenza viruses made employing reverse genetics techniques as described in US 5,166,057 and in Enami (1990) *Proc Natl Acad Sci* 87:3802-3805;
- 15 Enami & Palese (1991) *J Virol* 65:2711-2713 and Luytjes (1989) *Cell* 59:110, (see also McMichael (1983) *NEJ Med* 309:13, and Yap (1978) *Nature* 273:238 and *Nature* (1979) 277:108); human immunodeficiency virus as described in EP-0386882 and in Buchschacher (1992) *J. Virol.* 66:2731; measles virus, for example ATCC VR-67 and VR-1247 and those described in EP-0440219; Aura virus, for example ATCC VR-368; Bebaru virus, for example ATCC VR-600 and ATCC VR-1240;
- 20 Cabassou virus, for example ATCC VR-922; Chikungunya virus, for example ATCC VR-64 and ATCC VR-1241; Fort Morgan Virus, for example ATCC VR-924; Getah virus, for example ATCC VR-369 and ATCC VR-1243; Kyzylagach virus, for example ATCC VR-927; Mayaro virus, for example ATCC VR-66; Mucambo virus, for example ATCC VR-580 and ATCC VR-1244; Ndumu virus, for example ATCC VR-371; Pixuna virus, for example ATCC VR-372 and ATCC VR-1245;
- 25 Tonate virus, for example ATCC VR-925; Trinit virus, for example ATCC VR-469; Una virus, for example ATCC VR-374; Whataroa virus, for example ATCC VR-926; Y-62-33 virus, for example ATCC VR-375; O'Nyong virus, Eastern encephalitis virus, for example ATCC VR-65 and ATCC VR-1242; Western encephalitis virus, for example ATCC VR-70, ATCC VR-1251, ATCC VR-622 and ATCC VR-1252; and coronavirus, for example ATCC VR-740 and those described in Hamre
- 30 (1966) *Proc Soc Exp Biol Med* 121:190.

Delivery of the compositions of this invention into cells is not limited to the above mentioned viral vectors. Other delivery methods and media may be employed such as, for example, nucleic acid

expression vectors, polycationic condensed DNA linked or unlinked to killed adenovirus alone, for example see US Serial No. 08/366,787, filed December 30, 1994 and Curiel (1992) *Hum Gene Ther* 3:147-154 ligand linked DNA, for example see Wu (1989) *J Biol Chem* 264:16985-16987, eucaryotic cell delivery vehicles cells, for example see US Serial No.08/240,030, filed May 9, 5 1994, and US Serial No. 08/404,796, deposition of photopolymerized hydrogel materials, hand-held gene transfer particle gun, as described in US Patent 5,149,655, ionizing radiation as described in US5,206,152 and in WO92/11033, nucleic charge neutralization or fusion with cell membranes. Additional approaches are described in Philip (1994) *Mol Cell Biol* 14:2411-2418 and in Woffendin (1994) *Proc Natl Acad Sci* 91:1581-1585.

- 10 Particle mediated gene transfer may be employed, for example see US Serial No. 60/023,867. Briefly, the sequence can be inserted into conventional vectors that contain conventional control sequences for high level expression, and then incubated with synthetic gene transfer molecules such as polymeric DNA-binding cations like polylysine, protamine, and albumin, linked to cell targeting ligands such as asialoorosomucoid, as described in Wu & Wu (1987) *J. Biol. Chem.* 15 262:4429-4432, insulin as described in Hucked (1990) *Biochem Pharmacol* 40:253-263, galactose as described in Plank (1992) *Bioconjugate Chem* 3:533-539, lactose or transferrin.

Naked DNA may also be employed. Exemplary naked DNA introduction methods are described in WO 90/11092 and US 5,580,859. Uptake efficiency may be improved using biodegradable latex beads. DNA coated latex beads are efficiently transported into cells after endocytosis initiation by the 20 beads. The method may be improved further by treatment of the beads to increase hydrophobicity and thereby facilitate disruption of the endosome and release of the DNA into the cytoplasm.

Liposomes that can act as gene delivery vehicles are described in US 5,422,120, WO95/13796, WO94/23697, WO91/14445 and EP-524,968. As described in USSN. 60/023,867, on non-viral delivery, the nucleic acid sequences encoding a polypeptide can be inserted into conventional 25 vectors that contain conventional control sequences for high level expression, and then be incubated with synthetic gene transfer molecules such as polymeric DNA-binding cations like polylysine, protamine, and albumin, linked to cell targeting ligands such as asialoorosomucoid, insulin, galactose, lactose, or transferrin. Other delivery systems include the use of liposomes to encapsulate DNA comprising the gene under the control of a variety of tissue-specific or ubiquitously-active 30 promoters. Further non-viral delivery suitable for use includes mechanical delivery systems such as the approach described in Woffendin *et al* (1994) *Proc. Natl. Acad. Sci. USA*

91(24):11581-11585. Moreover, the coding sequence and the product of expression of such can be delivered through deposition of photopolymerized hydrogel materials. Other conventional methods for gene delivery that can be used for delivery of the coding sequence include, for example, use of hand-held gene transfer particle gun, as described in US 5,149,655; use of ionizing radiation for
5 activating transferred gene, as described in US 5,206,152 and WO92/11033

Exemplary liposome and polycationic gene delivery vehicles are those described in US 5,422,120 and 4,762,915; in WO 95/13796; WO94/23697; and WO91/14445; in EP-0524968; and in Stryer, Biochemistry, pages 236-240 (1975) W.H. Freeman, San Francisco; Szoka (1980) *Biochem Biophys Acta* 600:1; Bayer (1979) *Biochem Biophys Acta* 550:464; Rivnay (1987) *Meth Enzymol*
10 149:119; Wang (1987) *Proc Natl Acad Sci* 84:7851; Plant (1989) *Anal Biochem* 176:420.

A polynucleotide composition can comprises therapeutically effective amount of a gene therapy vehicle, as the term is defined above. For purposes of the present invention, an effective dose will be from about 0.01 mg/ kg to 50 mg/kg or 0.05 mg/kg to about 10 mg/kg of the DNA constructs in the individual to which it is administered.

15 Delivery Methods

Once formulated, the polynucleotide compositions of the invention can be administered (1) directly to the subject; (2) delivered *ex vivo*, to cells derived from the subject; or (3) *in vitro* for expression of recombinant proteins. The subjects to be treated can be mammals or birds. Also, human subjects can be treated.

20 Direct delivery of the compositions will generally be accomplished by injection, either subcutaneously, intraperitoneally, intravenously or intramuscularly or delivered to the interstitial space of a tissue. The compositions can also be administered into a lesion. Other modes of administration include oral and pulmonary administration, suppositories, and transdermal or transcutaneous applications (*eg.* see WO98/20734), needles, and gene guns or hyposprays. Dosage
25 treatment may be a single dose schedule or a multiple dose schedule.

Methods for the *ex vivo* delivery and reimplantation of transformed cells into a subject are known in the art and described in *eg.* WO93/14778. Examples of cells useful in *ex vivo* applications include, for example, stem cells, particularly hematopoietic, lymph cells, macrophages, dendritic cells, or tumor cells.

Generally, delivery of nucleic acids for both *ex vivo* and *in vitro* applications can be accomplished by the following procedures, for example, dextran-mediated transfection, calcium phosphate precipitation, polybrene mediated transfection, protoplast fusion, electroporation, encapsulation of the polynucleotide(s) in liposomes, and direct microinjection of the DNA into nuclei, all well known in the art.

Polynucleotide and polypeptide pharmaceutical compositions

In addition to the pharmaceutically acceptable carriers and salts described above, the following additional agents can be used with polynucleotide and/or polypeptide compositions.

A. Polypeptides

One example are polypeptides which include, without limitation: asialoglycoproteins; antibodies; antibody fragments; ferritin; interleukins; interferons, granulocyte, macrophage colony stimulating factor (GM-CSF), granulocyte colony stimulating factor (G-CSF), macrophage colony stimulating factor (M-CSF), stem cell factor and erythropoietin. Viral antigens, such as envelope proteins, can also be used. Also, proteins from other invasive organisms, such as the 17 amino acid peptide from the circumsporozoite protein of plasmodium falciparum known as RII.

B. Hormones, Vitamins, etc.

Other groups that can be included are, for example: hormones, steroids, androgens, estrogens, thyroid hormone, or vitamins, folic acid.

C. Polyalkylenes, Polysaccharides, etc.

Also, polyalkylene glycol can be included with the desired polynucleotides/polypeptides. In a preferred embodiment, the polyalkylene glycol is polyethylene glycol. In addition, mono-, di-, or polysaccharides can be included. In a preferred embodiment of this aspect, the polysaccharide is dextran or DEAE-dextran. Also, chitosan and poly(lactide-co-glycolide)

D. Lipids, and Liposomes

The desired polynucleotide/polypeptide can also be encapsulated in lipids or packaged in liposomes prior to delivery to the subject or to cells derived therefrom.

Lipid encapsulation is generally accomplished using liposomes which are able to stably bind or entrap and retain nucleic acid. The ratio of condensed polynucleotide to lipid preparation can vary but will generally be around 1:1 (mg DNA:micromoles lipid), or more of lipid. For a review of the

use of liposomes as carriers for delivery of nucleic acids, see, Hug and Sleight (1991) *Biochim. Biophys. Acta*. 1097:1-17; Straubinger (1983) *Meth. Enzymol.* 101:512-527.

Liposomal preparations for use in the present invention include cationic (positively charged), anionic (negatively charged) and neutral preparations. Cationic liposomes have been shown to
5 mediate intracellular delivery of plasmid DNA (Felgner (1987) *Proc. Natl. Acad. Sci. USA* 84:7413-7416); mRNA (Malone (1989) *Proc. Natl. Acad. Sci. USA* 86:6077-6081); and purified transcription factors (Debs (1990) *J. Biol. Chem.* 265:10189-10192), in functional form.

Cationic liposomes are readily available. For example, N[1-2,3-dioleoyloxy)propyl]-N,N,N-triethylammonium (DOTMA) liposomes are available under the trademark Lipofectin, from GIBCO BRL, Grand
10 Island, NY. (See, also, Felgner *supra*). Other commercially available liposomes include transfectace (DDAB/DOPE) and DOTAP/DOPE (Boehringer). Other cationic liposomes can be prepared from readily available materials using techniques well known in the art. See, *eg.* Szoka (1978) *Proc. Natl. Acad. Sci. USA* 75:4194-4198; WO90/11092 for a description of the synthesis of DOTAP (1,2-bis(oleoyloxy)-3-(trimethylammonio)propane) liposomes.

15 Similarly, anionic and neutral liposomes are readily available, such as from Avanti Polar Lipids (Birmingham, AL), or can be easily prepared using readily available materials. Such materials include phosphatidyl choline, cholesterol, phosphatidyl ethanolamine, dioleoylphosphatidyl choline (DOPC), dioleoylphosphatidyl glycerol (DOPG), dioleoylphosphatidyl ethanolamine (DOPE), among others. These materials can also be mixed with the DOTMA and DOTAP starting materials in appropriate
20 ratios. Methods for making liposomes using these materials are well known in the art.

The liposomes can comprise multilammellar vesicles (MLVs), small unilamellar vesicles (SUVs), or large unilamellar vesicles (LUVs). The various liposome-nucleic acid complexes are prepared using methods known in the art. See *eg.* Straubinger (1983) *Meth. Immunol.* 101:512-527; Szoka (1978) *Proc. Natl. Acad. Sci. USA* 75:4194-4198; Papahadjopoulos (1975) *Biochim. Biophys. Acta*
25 394:483; Wilson (1979) *Cell* 17:77; Deamer & Bangham (1976) *Biochim. Biophys. Acta* 443:629; Ostro (1977) *Biochem. Biophys. Res. Commun.* 76:836; Fraley (1979) *Proc. Natl. Acad. Sci. USA* 76:3348; Enoch & Strittmatter (1979) *Proc. Natl. Acad. Sci. USA* 76:145; Fraley (1980) *J. Biol. Chem.* (1980) 255:10431; Szoka & Papahadjopoulos (1978) *Proc. Natl. Acad. Sci. USA* 75:145; and Schaefer-Ridder (1982) *Science* 215:166.

E. Lipoproteins

In addition, lipoproteins can be included with the polynucleotide/polypeptide to be delivered. Examples of lipoproteins to be utilized include: chylomicrons, HDL, IDL, LDL, and VLDL. Mutants, fragments, or fusions of these proteins can also be used. Also, modifications of naturally occurring lipoproteins can be used, such as acetylated LDL. These lipoproteins can target the delivery of polynucleotides to cells expressing lipoprotein receptors. Preferably, if lipoproteins are including with the polynucleotide to be delivered, no other targeting ligand is included in the composition.

Naturally occurring lipoproteins comprise a lipid and a protein portion. The protein portion are known as apoproteins. At the present, apoproteins A, B, C, D, and E have been isolated and identified. At least two of these contain several proteins, designated by Roman numerals, AI, AII, AIV; CI, CII, CIII.

A lipoprotein can comprise more than one apoprotein. For example, naturally occurring chylomicrons comprises of A, B, C, and E, over time these lipoproteins lose A and acquire C and E apoproteins. VLDL comprises A, B, C, and E apoproteins, LDL comprises apoprotein B; and HDL comprises apoproteins A, C, and E.

The amino acid of these apoproteins are known and are described in, for example, Breslow (1985) *Annu Rev. Biochem* 54:699; Law (1986) *Adv. Exp Med. Biol.* 151:162; Chen (1986) *J Biol Chem* 261:12918; Kane (1980) *Proc Natl Acad Sci USA* 77:2465; and Utermann (1984) *Hum Genet* 65:232.

Lipoproteins contain a variety of lipids including, triglycerides, cholesterol (free and esters), and phospholipids. The composition of the lipids varies in naturally occurring lipoproteins. For example, chylomicrons comprise mainly triglycerides. A more detailed description of the lipid content of naturally occurring lipoproteins can be found, for example, in *Meth. Enzymol.* 128 (1986). The composition of the lipids are chosen to aid in conformation of the apoprotein for receptor binding activity. The composition of lipids can also be chosen to facilitate hydrophobic interaction and association with the polynucleotide binding molecule.

Naturally occurring lipoproteins can be isolated from serum by ultracentrifugation, for instance. Such methods are described in *Meth. Enzymol. (supra)*; Pitas (1980) *J. Biochem.* 255:5454-5460 and Mahey (1979) *J Clin. Invest* 64:743-750. Lipoproteins can also be produced by *in vitro* or recombinant methods by expression of the apoprotein genes in a desired host cell. See, for example, Atkinson (1986) *Annu Rev Biophys Chem* 15:403 and Radding (1958) *Biochim Biophys Acta* 30:

443. Lipoproteins can also be purchased from commercial suppliers, such as Biomedical Technologies, Inc., Stoughton, Massachusetts, USA. Further description of lipoproteins can be found in Zuckermann *et al.* PCT/US97/14465.

F. Polycationic Agents

- 5 Polycationic agents can be included, with or without lipoprotein, in a composition with the desired polynucleotide/polypeptide to be delivered.

Polycationic agents, typically, exhibit a net positive charge at physiological relevant pH and are capable of neutralizing the electrical charge of nucleic acids to facilitate delivery to a desired location. These agents have both in vitro, ex vivo, and in vivo applications. Polycationic agents can
10 be used to deliver nucleic acids to a living subject either intramuscularly, subcutaneously, etc.

The following are examples of useful polypeptides as polycationic agents: polylysine, polyarginine, polyornithine, and protamine. Other examples include histones, protamines, human serum albumin, DNA binding proteins, non-histone chromosomal proteins, coat proteins from DNA viruses, such as (X174, transcriptional factors also contain domains that bind DNA and therefore may be useful
15 as nucleic acid condensing agents. Briefly, transcriptional factors such as C/CEBP, c-jun, c-fos, AP-1, AP-2, AP-3, CPF, Prot-1, Sp-1, Oct-1, Oct-2, CREP, and TFIID contain basic domains that bind DNA sequences.

Organic polycationic agents include: spermine, spermidine, and putrescine.

The dimensions and of the physical properties of a polycationic agent can be extrapolated from the
20 list above, to construct other polypeptide polycationic agents or to produce synthetic polycationic agents.

Synthetic polycationic agents which are useful include, for example, DEAE-dextran, polybrene. Lipofectin™, and lipofectAMINE™ are monomers that form polycationic complexes when combined with polynucleotides/polypeptides.

25 Immunodiagnostic Assays

Neisserial antigens of the invention can be used in immunoassays to detect antibody levels (or, conversely, anti-Neisserial antibodies can be used to detect antigen levels). Immunoassays based on well defined, recombinant antigens can be developed to replace invasive diagnostics methods. Antibodies to Neisserial proteins within biological samples, including for example, blood or serum

samples, can be detected. Design of the immunoassays is subject to a great deal of variation, and a variety of these are known in the art. Protocols for the immunoassay may be based, for example, upon competition, or direct reaction, or sandwich type assays. Protocols may also, for example, use solid supports, or may be by immunoprecipitation. Most assays involve the use of labeled antibody
5 or polypeptide; the labels may be, for example, fluorescent, chemiluminescent, radioactive, or dye molecules. Assays which amplify the signals from the probe are also known; examples of which are assays which utilize biotin and avidin, and enzyme-labeled and mediated immunoassays, such as ELISA assays.

Kits suitable for immunodiagnosis and containing the appropriate labeled reagents are constructed
10 by packaging the appropriate materials, including the compositions of the invention, in suitable containers, along with the remaining reagents and materials (for example, suitable buffers, salt solutions, *etc.*) required for the conduct of the assay, as well as suitable set of assay instructions.

Nucleic Acid Hybridisation

“Hybridization” refers to the association of two nucleic acid sequences to one another by hydrogen
15 bonding. Typically, one sequence will be fixed to a solid support and the other will be free in solution. Then, the two sequences will be placed in contact with one another under conditions that favor hydrogen bonding. Factors that affect this bonding include: the type and volume of solvent; reaction temperature; time of hybridization; agitation; agents to block the non-specific attachment of the liquid phase sequence to the solid support (Denhardt's reagent or BLOTTO); concentration of the sequences;
20 use of compounds to increase the rate of association of sequences (dextran sulfate or polyethylene glycol); and the stringency of the washing conditions following hybridization. See Sambrook *et al.* [*supra*] Volume 2, chapter 9, pages 9.47 to 9.57.

“Stringency” refers to conditions in a hybridization reaction that favor association of very similar sequences over sequences that differ. For example, the combination of temperature and salt
25 concentration should be chosen that is approximately 120 to 200°C below the calculated T_m of the hybrid under study. The temperature and salt conditions can often be determined empirically in preliminary experiments in which samples of genomic DNA immobilized on filters are hybridized to the sequence of interest and then washed under conditions of different stringencies. See Sambrook *et al.* at page 9.50.

30 Variables to consider when performing, for example, a Southern blot are (1) the complexity of the DNA being blotted and (2) the homology between the probe and the sequences being detected. The

- total amount of the fragment(s) to be studied can vary a magnitude of 10, from 0.1 to 1 µg for a plasmid or phage digest to 10^{-9} to 10^{-8} g for a single copy gene in a highly complex eukaryotic genome. For lower complexity polynucleotides, substantially shorter blotting, hybridization, and exposure times, a smaller amount of starting polynucleotides, and lower specific activity of probes can be used. For example, a single-copy yeast gene can be detected with an exposure time of only 1 hour starting with 1 µg of yeast DNA, blotting for two hours, and hybridizing for 4-8 hours with a probe of 10^8 cpm/µg. For a single-copy mammalian gene a conservative approach would start with 10 µg of DNA, blot overnight, and hybridize overnight in the presence of 10% dextran sulfate using a probe of greater than 10^8 cpm/µg, resulting in an exposure time of ~24 hours.
- Several factors can affect the melting temperature (T_m) of a DNA-DNA hybrid between the probe and the fragment of interest, and consequently, the appropriate conditions for hybridization and washing. In many cases the probe is not 100% homologous to the fragment. Other commonly encountered variables include the length and total G+C content of the hybridizing sequences and the ionic strength and formamide content of the hybridization buffer. The effects of all of these factors can be approximated by a single equation:

$$T_m = 81 + 16.6(\log_{10} C_i) + 0.4[\%(G + C)] - 0.6(\% \text{formamide}) - 600/n - 1.5(\% \text{mismatch}).$$

where C_i is the salt concentration (monovalent ions) and n is the length of the hybrid in base pairs (slightly modified from Meinkoth & Wahl (1984) *Anal. Biochem.* 138: 267-284).

- In designing a hybridization experiment, some factors affecting nucleic acid hybridization can be conveniently altered. The temperature of the hybridization and washes and the salt concentration during the washes are the simplest to adjust. As the temperature of the hybridization increases (*ie.* stringency), it becomes less likely for hybridization to occur between strands that are nonhomologous, and as a result, background decreases. If the radiolabeled probe is not completely homologous with the immobilized fragment (as is frequently the case in gene family and interspecies hybridization experiments), the hybridization temperature must be reduced, and background will increase. The temperature of the washes affects the intensity of the hybridizing band and the degree of background in a similar manner. The stringency of the washes is also increased with decreasing salt concentrations.

- In general, convenient hybridization temperatures in the presence of 50% formamide are 42°C for a probe with is 95% to 100% homologous to the target fragment, 37°C for 90% to 95% homology,

and 32°C for 85% to 90% homology. For lower homologies, formamide content should be lowered and temperature adjusted accordingly, using the equation above. If the homology between the probe and the target fragment are not known, the simplest approach is to start with both hybridization and wash conditions which are nonstringent. If non-specific bands or high background are observed
5 after autoradiography, the filter can be washed at high stringency and reexposed. If the time required for exposure makes this approach impractical, several hybridization and/or washing stringencies should be tested in parallel.

Nucleic Acid Probe Assays

Methods such as PCR, branched DNA probe assays, or blotting techniques utilizing nucleic acid
10 probes according to the invention can determine the presence of cDNA or mRNA. A probe is said to "hybridize" with a sequence of the invention if it can form a duplex or double stranded complex, which is stable enough to be detected.

The nucleic acid probes will hybridize to the Neisserial nucleotide sequences of the invention (including both sense and antisense strands). Though many different nucleotide sequences will
15 encode the amino acid sequence, the native Neisserial sequence is preferred because it is the actual sequence present in cells. mRNA represents a coding sequence and so a probe should be complementary to the coding sequence; single-stranded cDNA is complementary to mRNA, and so a cDNA probe should be complementary to the non-coding sequence.

The probe sequence need not be identical to the Neisserial sequence (or its complement) — some
20 variation in the sequence and length can lead to increased assay sensitivity if the nucleic acid probe can form a duplex with target nucleotides, which can be detected. Also, the nucleic acid probe can include additional nucleotides to stabilize the formed duplex. Additional Neisserial sequence may also be helpful as a label to detect the formed duplex. For example, a non-complementary nucleotide sequence may be attached to the 5' end of the probe, with the remainder of the probe
25 sequence being complementary to a Neisserial sequence. Alternatively, non-complementary bases or longer sequences can be interspersed into the probe, provided that the probe sequence has sufficient complementarity with the a Neisserial sequence in order to hybridize therewith and thereby form a duplex which can be detected.

The exact length and sequence of the probe will depend on the hybridization conditions, such as
30 temperature, salt condition and the like. For example, for diagnostic applications, depending on the

complexity of the analyte sequence, the nucleic acid probe typically contains at least 10-20 nucleotides, preferably 15-25, and more preferably at least 30 nucleotides, although it may be shorter than this. Short primers generally require cooler temperatures to form sufficiently stable hybrid complexes with the template.

- 5 Probes may be produced by synthetic procedures, such as the triester method of Matteucci *et al.* [*J. Am. Chem. Soc.* (1981) 103:3185], or according to Urdea *et al.* [*Proc. Natl. Acad. Sci. USA* (1983) 80: 7461], or using commercially available automated oligonucleotide synthesizers.

The chemical nature of the probe can be selected according to preference. For certain applications, DNA or RNA are appropriate. For other applications, modifications may be incorporated *eg.*
10 backbone modifications, such as phosphorothioates or methylphosphonates, can be used to increase *in vivo* half-life, alter RNA affinity, increase nuclease resistance *etc.* [*eg.* see Agrawal & Iyer (1995) *Curr Opin Biotechnol* 6:12-19; Agrawal (1996) *TIBTECH* 14:376-387]; analogues such as peptide nucleic acids may also be used [*eg.* see Corey (1997) *TIBTECH* 15:224-229; Buchardt *et al.* (1993) *TIBTECH* 11:384-386].

- 15 Alternatively, the polymerase chain reaction (PCR) is another well-known means for detecting small amounts of target nucleic acids. The assay is described in: Mullis *et al.* [*Meth. Enzymol.* (1987) 155: 335-350]; US patents 4,683,195 and 4,683,202. Two "primer" nucleotides hybridize with the target nucleic acids and are used to prime the reaction. The primers can comprise sequence that does not hybridize to the sequence of the amplification target (or its complement) to aid with
20 duplex stability or, for example, to incorporate a convenient restriction site. Typically, such sequence will flank the desired Neisserial sequence.

A thermostable polymerase creates copies of target nucleic acids from the primers using the original target nucleic acids as a template. After a threshold amount of target nucleic acids are generated by the polymerase, they can be detected by more traditional methods, such as Southern
25 blots. When using the Southern blot method, the labelled probe will hybridize to the Neisserial sequence (or its complement).

Also, mRNA or cDNA can be detected by traditional blotting techniques described in Sambrook *et al* [*supra*]. mRNA, or cDNA generated from mRNA using a polymerase enzyme, can be purified and separated using gel electrophoresis. The nucleic acids on the gel are then blotted onto a solid
30 support, such as nitrocellulose. The solid support is exposed to a labelled probe and then washed

to remove any unhybridized probe. Next, the duplexes containing the labeled probe are detected. Typically, the probe is labelled with a radioactive moiety.

BRIEF DESCRIPTION OF THE DRAWINGS

Figures 1-20 show biochemical data obtained in the Examples, and also sequence analysis, for ORFs 37, 5, 2, 15, 22, 28, 32, 4, 61, 76, 89, 97, 106, 138, 23, 25, 27, 79, 85 and 132. M1 and M2 are molecular weight markers. Arrows indicate the position of the main recombinant product or, in Western blots, the position of the main *N.meningitidis* immunoreactive band. TP indicates *N.meningitidis* total protein extract; OMV indicates *N.meningitidis* outer membrane vesicle preparation. In bactericidal assay results: a diamond (◆) shows preimmune data; a triangle (▲) shows GST control data; a circle (●) shows data with recombinant *N.meningitidis* protein. Computer analyses show a hydrophilicity plot (upper), an antigenic index plot (middle), and an AMPHI analysis (lower). The AMPHI program has been used to predict T-cell epitopes [Gao *et al.* (1989) *J. Immunol.* **143**:3007; Roberts *et al.* (1996) *AIDS Res Hum Retrovir* **12**:593; Quakyi *et al.* (1992) *Scand J Immunol suppl.* **11**:9) and is available in the Protean package of DNASTAR, Inc. (1228 South Park Street, Madison, Wisconsin 53715 USA).

EXAMPLES

The examples describe nucleic acid sequences which have been identified in *N.meningitidis*, along with their putative translation products, and also those of *N.gonorrhoeae*. Not all of the nucleic acid sequences are complete *ie.* they encode less than the full-length wild-type protein.

The examples are generally in the following format:

- a nucleotide sequence which has been identified in *N.meningitidis* (strain B)
- the putative translation product of this sequence
- a computer analysis of the translation product based on database comparisons
- corresponding gene and protein sequences identified in *N.meningitidis* (strain A) and in *N.gonorrhoeae*
- a description of the characteristics of the proteins which indicates that they might be suitably antigenic
- results of biochemical analysis (expression, purification, ELISA, FACS *etc.*)

The examples typically include details of sequence identity between species and strains. Proteins that are similar in sequence are generally similar in both structure and function, and the sequence identity often indicates a common evolutionary origin. Comparison with sequences of proteins of known function is widely used as a guide for the assignment of putative protein function to a new sequence and has proved particularly useful in whole-genome analyses.

Sequence comparisons were performed at NCBI (<http://www.ncbi.nlm.nih.gov>) using the algorithms BLAST, BLAST2, BLASTn, BLASTp, tBLASTn, BLASTx, & tBLASTx [eg. see also Altschul *et al.* (1997) Gapped BLAST and PSI-BLAST: a new generation of protein database search programs. *Nucleic Acids Research* 25:2289-3402]. Searches were performed against the following databases: non-redundant GenBank+EMBL+DDBJ+PDB sequences and non-redundant GenBank CDS translations+PDB+SwissProt+SPupdate+PIR sequences.

To compare Meningococcal and Gonococcal sequences, the tBLASTx algorithm was used, as implemented at http://www.genome.ou.edu/gono_blast.html. The FASTA algorithm was also used to compare the ORFs (from GCG Wisconsin Package, version 9.0).

Dots within nucleotide sequences (eg. position 495 in SEQ ID 11) represent nucleotides which have been arbitrarily introduced in order to maintain a reading frame. In the same way, double-underlined nucleotides were removed. Lower case letters (eg. position 496 in SEQ ID 11) represent ambiguities which arose during alignment of independent sequencing reactions (some of the nucleotide sequences in the examples are derived from combining the results of two or more experiments).

Nucleotide sequences were scanned in all six reading frames to predict the presence of hydrophobic domains using an algorithm based on the statistical studies of Esposti *et al.* [Critical evaluation of the hydropathy of membrane proteins (1990) *Eur J Biochem* 190:207-219]. These domains represent potential transmembrane regions or hydrophobic leader sequences.

Open reading frames were predicted from fragmented nucleotide sequences using the program ORFFINDER (NCBI).

Underlined amino acid sequences indicate possible transmembrane domains or leader sequences in the ORFs, as predicted by the PSORT algorithm (<http://www.psорт.nibb.ac.jp>). Functional domains were also predicted using the MOTIFS program (GCG Wisconsin & PROSITE).

Various tests can be used to assess the *in vivo* immunogenicity of the proteins identified in the examples. For example, the proteins can be expressed recombinantly and used to screen patient sera by immunoblot. A positive reaction between the protein and patient serum indicates that the patient has previously mounted an immune response to the protein in question *ie.* the protein is an immunogen. This method can also be used to identify immunodominant proteins.

The recombinant protein can also be conveniently used to prepare antibodies *eg.* in a mouse. These can be used for direct confirmation that a protein is located on the cell-surface. Labelled antibody (*eg.* fluorescent labelling for FACS) can be incubated with intact bacteria and the presence of label on the bacterial surface confirms the location of the protein.

10 In particular, the following methods (A) to (S) were used to express, purify and biochemically characterise the proteins of the invention:

A) Chromosomal DNA preparation

N.meningitidis strain 2996 was grown to exponential phase in 100ml of GC medium, harvested by centrifugation, and resuspended in 5ml buffer (20% Sucrose, 50mM Tris-HCl, 50mM EDTA, pH8).

15 After 10 minutes incubation on ice, the bacteria were lysed by adding 10ml lysis solution (50mM NaCl, 1% Na-Sarkosyl, 50µg/ml Proteinase K), and the suspension was incubated at 37°C for 2 hours. Two phenol extractions (equilibrated to pH 8) and one ChCl_3 /isoamylalcohol (24:1) extraction were performed. DNA was precipitated by addition of 0.3M sodium acetate and 2 volumes ethanol, and was collected by centrifugation. The pellet was washed once with 70%

20 ethanol and redissolved in 4ml buffer (10mM Tris-HCl, 1mM EDTA, pH 8). The DNA concentration was measured by reading the OD at 260 nm.

B) Oligonucleotide design

Synthetic oligonucleotide primers were designed on the basis of the coding sequence of each ORF, using (a) the meningococcus B sequence when available, or (b) the gonococcus/meningococcus A

25 sequence, adapted to the codon preference usage of meningococcus as necessary. Any predicted signal peptides were omitted, by deducing the 5'-end amplification primer sequence immediately downstream from the predicted leader sequence.

For most ORFs, the 5' primers included two restriction enzyme recognition sites (*Bam*HI-*Nde*I, *Bam*HI-*Nhe*I, or *Eco*RI-*Nhe*I, depending on the gene's own restriction pattern); the 3' primers included

a *XhoI* restriction site. This procedure was established in order to direct the cloning of each amplification product (corresponding to each ORF) into two different expression systems: pGEX-KG (using either *BamHI-XhoI* or *EcoRI-XhoI*), and pET21b+ (using either *NdeI-XhoI* or *NheI-XhoI*).

5 5'-end primer tail: CGCGGATCCCATATG (*BamHI-NdeI*)
 CGCGGATCCGCTAGC (*BamHI-NheI*)
 CCGGAATTCTAGCTAGC (*EcoRI-NheI*)
 3'-end primer tail: CCCGCTCGAG (*XhoI*)

For ORFs 5, 15, 17, 19, 20, 22, 27, 28, 65 & 89, two different amplifications were performed to clone each ORF in the two expression systems. Two different 5' primers were used for each ORF;
 10 the same 3' *XhoI* primer was used as before:

5'-end primer tail: GGAATTCATATGGCCATGG (*NdeI*)
 5'-end primer tail: CGGGATCC (*BamHI*)

ORF 76 was cloned in the pTRC expression vector and expressed as an amino-terminus His-tag fusion. In this particular case, the predicted signal peptide was included in the final product. *NheI*-
 15 *BamHI* restriction sites were incorporated using primers:

5'-end primer tail: GATCAGCTAGCCATATG (*NheI*)
 3'-end primer tail: CGGGATCC (*BamHI*)

As well as containing the restriction enzyme recognition sequences, the primers included nucleotides which hybridized to the sequence to be amplified. The number of hybridizing
 20 nucleotides depended on the melting temperature of the whole primer, and was determined for each primer using the formulae:

$$T_m = 4 (G+C) + 2 (A+T) \quad (\text{tail excluded})$$

$$T_m = 64.9 + 0.41 (\% \text{ GC}) - 600/N \quad (\text{whole primer})$$

The average melting temperature of the selected oligos were 65-70°C for the whole oligo and
 25 50-55°C for the hybridising region alone.

Table I (page 487) shows the forward and reverse primers used for each amplification. In certain cases, it will be noted that the sequence of the primer does not exactly match the sequence in the ORF. When initial amplifications were performed, the complete 5' and/or 3' sequence was not

known for some meningococcal ORFs, although the corresponding sequences had been identified in gonococcus. For amplification, the gonococcal sequences could thus be used as the basis for primer design, altered to take account of codon preference. In particular, the following codons were changed: ATA→ATT; TCG→TCT; CAG→CAA; AAG→AAA; GAG→GAA; CGA→CGC;
 5 CGG→CGC; GGG→GGC. Italicised nucleotides in Table I indicate such a change. It will be appreciated that, once the complete sequence has been identified, this approach is generally no longer necessary.

Oligos were synthesized by a Perkin Elmer 394 DNA/RNA Synthesizer, eluted from the columns in 2ml NH₄OH, and deprotected by 5 hours incubation at 56°C. The oligos were precipitated by
 10 addition of 0.3M Na-Acetate and 2 volumes ethanol. The samples were then centrifuged and the pellets resuspended in either 100µl or 1ml of water. OD₂₆₀ was determined using a Perkin Elmer Lambda Bio spectrophotometer and the concentration was determined and adjusted to 2-10pmol/µl.

C) Amplification

The standard PCR protocol was as follows: 50-200ng of genomic DNA were used as a template
 15 in the presence of 20-40µM of each oligo, 400-800µM dNTPs solution, 1x PCR buffer (including 1.5mM MgCl₂), 2.5 units *TaqI* DNA polymerase (using Perkin-Elmer AmpliTaq, GIBCO Platinum, Pwo DNA polymerase, or Tahara Shuzo Taq polymerase).

In some cases, PCR was optimised by the addition of 10µl DMSO or 50µl 2M betaine.

After a hot start (adding the polymerase during a preliminary 3 minute incubation of the whole mix
 20 at 95°C), each sample underwent a double-step amplification: the first 5 cycles were performed using as the hybridization temperature the one of the oligos excluding the restriction enzymes tail, followed by 30 cycles performed according to the hybridization temperature of the whole length oligos. The cycles were followed by a final 10 minute extension step at 72°C.

The standard cycles were as follows:

	Denaturation	Hybridisation	Elongation
First 5 cycles	30 seconds 95°C	30 seconds 50-55°C	30-60 seconds 72°C
Last 30 cycles	30 seconds	30 seconds	30-60 seconds

	95°C	65-70°C	72°C
--	------	---------	------

The elongation time varied according to the length of the ORF to be amplified.

The amplifications were performed using either a 9600 or a 2400 Perkin Elmer GeneAmp PCR System. To check the results, 1/10 of the amplification volume was loaded onto a 1-1.5% agarose gel and the size of each amplified fragment compared with a DNA molecular weight marker.

- 5 The amplified DNA was either loaded directly on a 1% agarose gel or first precipitated with ethanol and resuspended in a suitable volume to be loaded on a 1% agarose gel. The DNA fragment corresponding to the right size band was then eluted and purified from gel, using the Qiagen Gel Extraction Kit, following the instructions of the manufacturer. The final volume of the DNA fragment was 30µl or 50µl of either water or 10mM Tris, pH 8.5.

10 D) Digestion of PCR fragments

The purified DNA corresponding to the amplified fragment was split into 2 aliquots and double-digested with:

- *NdeI/XhoI* or *NheI/XhoI* for cloning into pET-21b+ and further expression of the protein as a C-terminus His-tag fusion
- 15 – *BamHI/XhoI* or *EcoRI/XhoI* for cloning into pGEX-KG and further expression of the protein as N-terminus GST fusion.
- For ORF 76, *NheI/BamHI* for cloning into pTRC-HisA vector and further expression of the protein as N-terminus His-tag fusion.
- *EcoRI/PstI*, *EcoRI/SalI*, *SalI/PstI* for cloning into pGex-His and further expression of
- 20 the protein as N-terminus His-tag fusion

- Each purified DNA fragment was incubated (37°C for 3 hours to overnight) with 20 units of each restriction enzyme (New England Biolabs) in a either 30 or 40µl final volume in the presence of the appropriate buffer. The digestion product was then purified using the QIAquick PCR purification kit, following the manufacturer's instructions, and eluted in a final volume of 30 or
- 25 50µl of either water or 10mM Tris-HCl, pH 8.5. The final DNA concentration was determined by 1% agarose gel electrophoresis in the presence of titrated molecular weight marker.

E) Digestion of the cloning vectors (pET22B, pGEX-KG, pTRC-His A, and pGex-His)

10µg plasmid was double-digested with 50 units of each restriction enzyme in 200µl reaction volume in the presence of appropriate buffer by overnight incubation at 37°C. After loading the whole digestion on a 1% agarose gel, the band corresponding to the digested vector was purified
5 from the gel using the Qiagen QIAquick Gel Extraction Kit and the DNA was eluted in 50µl of 10mM Tris-HCl, pH 8.5. The DNA concentration was evaluated by measuring OD₂₆₀ of the sample, and adjusted to 50µg/µl. 1µl of plasmid was used for each cloning procedure.

The vector pGEX-His is a modified pGEX-2T vector carrying a region encoding six histidine residues upstream to the thrombin cleavage site and containing the multiple cloning site of the
10 vector pTRC99 (Pharmacia).

F) Cloning

The fragments corresponding to each ORF, previously digested and purified, were ligated in both pET22b and pGEX-KG. In a final volume of 20µl, a molar ratio of 3:1 fragment/vector was ligated using 0.5µl of NEB T4 DNA ligase (400 units/µl), in the presence of the buffer supplied by the manufacturer.
15 The reaction was incubated at room temperature for 3 hours. In some experiments, ligation was performed using the Boehringer "Rapid Ligation Kit", following the manufacturer's instructions.

In order to introduce the recombinant plasmid in a suitable strain, 100µl *E. coli* DH5 competent cells were incubated with the ligase reaction solution for 40 minutes on ice, then at 37°C for 3 minutes, then, after adding 800µl LB broth, again at 37°C for 20 minutes. The cells were then
20 centrifuged at maximum speed in an Eppendorf microfuge and resuspended in approximately 200µl of the supernatant. The suspension was then plated on LB ampicillin (100mg/ml).

The screening of the recombinant clones was performed by growing 5 randomly-chosen colonies overnight at 37°C in either 2ml (pGEX or pTC clones) or 5ml (pET clones) LB broth + 100µg/ml ampicillin. The cells were then pelleted and the DNA extracted using the Qiagen QIAprep Spin
25 Miniprep Kit, following the manufacturer's instructions, to a final volume of 30µl. 5µl of each individual miniprep (approximately 1g) were digested with either *NdeI/XhoI* or *BamHI/XhoI* and the whole digestion loaded onto a 1-1.5% agarose gel (depending on the expected insert size), in parallel with the molecular weight marker (1Kb DNA Ladder, GIBCO). The screening of the positive clones was made on the base of the correct insert size.

For the cloning of ORFs 110, 111, 113, 115, 119, 122, 125 & 130, the double-digested PCR product was ligated into double-digested vector using *EcoRI-PstI* cloning sites or, for ORFs 115 & 127, *EcoRI-SalI* or, for ORF 122, *SalI-PstI*. After cloning, the recombinant plasmids were introduced in the *E.coli* host W3110. Individual clones were grown overnight at 37°C in L-broth with 50µl/ml ampicillin.

G) Expression

Each ORF cloned into the expression vector was transformed into the strain suitable for expression of the recombinant protein product. 1µl of each construct was used to transform 30µl of *E.coli* BL21 (pGEX vector), *E.coli* TOP 10 (pTRC vector) or *E.coli* BL21-DE3 (pET vector), as described above. In the case of the pGEX-His vector, the same *E.coli* strain (W3110) was used for initial cloning and expression. Single recombinant colonies were inoculated into 2ml LB+Amp (100µg/ml), incubated at 37°C overnight, then diluted 1:30 in 20ml of LB+Amp (100µg/ml) in 100ml flasks, making sure that the OD₆₀₀ ranged between 0.1 and 0.15. The flasks were incubated at 30°C into gyratory water bath shakers until OD indicated exponential growth suitable for induction of expression (0.4-0.8 OD for pET and pTRC vectors; 0.8-1 OD for pGEX and pGEX-His vectors). For the pET, pTRC and pGEX-His vectors, the protein expression was induced by addition of 1mM IPTG, whereas in the case of pGEX system the final concentration of IPTG was 0.2mM. After 3 hours incubation at 30°C, the final concentration of the sample was checked by OD. In order to check expression, 1ml of each sample was removed, centrifuged in a microfuge, the pellet resuspended in PBS, and analysed by 12% SDS-PAGE with Coomassie Blue staining. The whole sample was centrifuged at 6000g and the pellet resuspended in PBS for further use.

H) GST-fusion proteins large-scale purification.

A single colony was grown overnight at 37°C on LB+Amp agar plate. The bacteria were inoculated into 20ml of LB+Amp liquid culture in a water bath shaker and grown overnight. Bacteria were diluted 1:30 into 600ml of fresh medium and allowed to grow at the optimal temperature (20-37°C) to OD₅₅₀ 0.8-1. Protein expression was induced with 0.2mM IPTG followed by three hours incubation. The culture was centrifuged at 8000rpm at 4°C. The supernatant was discarded and the bacterial pellet was resuspended in 7.5ml cold PBS. The cells were disrupted by sonication on ice for 30 sec at 40W using a Branson sonifier B-15, frozen and thawed twice and centrifuged again. The supernatant was collected and mixed with 150µl Glutathione-Sepharose 4B resin (Pharmacia)

(previously washed with PBS) and incubated at room temperature for 30 minutes. The sample was centrifuged at 700g for 5 minutes at 4°C. The resin was washed twice with 10ml cold PBS for 10 minutes, resuspended in 1ml cold PBS, and loaded on a disposable column. The resin was washed twice with 2ml cold PBS until the flow-through reached OD₂₈₀ of 0.02-0.06. The GST-fusion protein was eluted by addition of 700µl cold Glutathione elution buffer (10mM reduced glutathione, 50mM Tris-HCl) and fractions collected until the OD₂₈₀ was 0.1. 21µl of each fraction were loaded on a 12% SDS gel using either Biorad SDS-PAGE Molecular weight standard broad range (M1) (200, 116.25, 97.4, 66.2, 45, 31, 21.5, 14.4, 6.5 kDa) or Amersham Rainbow Marker (M2) (220, 66, 46, 30, 21.5, 14.3 kDa) as standards. As the MW of GST is 26kDa, this value must be added to the MW of each GST-fusion protein.

I) His-fusion solubility analysis (ORFs 111-129)

To analyse the solubility of the His-fusion expression products, pellets of 3ml cultures were resuspended in buffer M1 [500µl PBS pH 7.2]. 25µl lysozyme (10mg/ml) was added and the bacteria were incubated for 15 min at 4°C. The pellets were sonicated for 30 sec at 40W using a Branson sonifier B-15, frozen and thawed twice and then separated again into pellet and supernatant by a centrifugation step. The supernatant was collected and the pellet was resuspended in buffer M2 [8M urea, 0.5M NaCl, 20mM imidazole and 0.1M NaH₂PO₄] and incubated for 3 to 4 hours at 4°C. After centrifugation, the supernatant was collected and the pellet was resuspended in buffer M3 [6M guanidinium-HCl, 0.5M NaCl, 20mM imidazole and 0.1M NaH₂PO₄] overnight at 4°C. The supernatants from all steps were analysed by SDS-PAGE.

The proteins expressed from ORFs 113, 119 and 120 were found to be soluble in PBS, whereas ORFs 111, 122, 126 and 129 need urea and ORFs 125 and 127 need guanidinium-HCl for their solubilization.

J) His-fusion large-scale purification.

A single colony was grown overnight at 37°C on a LB + Amp agar plate. The bacteria were inoculated into 20ml of LB+Amp liquid culture and incubated overnight in a water bath shaker. Bacteria were diluted 1:30 into 600ml fresh medium and allowed to grow at the optimal temperature (20-37°C) to OD₅₅₀ 0.6-0.8. Protein expression was induced by addition of 1mM IPTG and the culture further incubated for three hours. The culture was centrifuged at 8000rpm at 4°C, the supernatant was discarded and the bacterial pellet was resuspended in 7.5ml of either (i) cold

buffer A (300mM NaCl, 50mM phosphate buffer, 10mM imidazole, pH 8) for soluble proteins or (ii) buffer B (urea 8M, 10mM Tris-HCl, 100mM phosphate buffer, pH 8.8) for insoluble proteins.

The cells were disrupted by sonication on ice for 30 sec at 40W using a Branson sonifier B-15, frozen and thawed two times and centrifuged again.

- 5 For insoluble proteins, the supernatant was stored at -20°C, while the pellets were resuspended in 2ml buffer C (6M guanidine hydrochloride, 100mM phosphate buffer, 10mM Tris-HCl, pH 7.5) and treated in a homogenizer for 10 cycles. The product was centrifuged at 13000rpm for 40 minutes.

Supernatants were collected and mixed with 150µl Ni²⁺-resin (Pharmacia) (previously washed with either buffer A or buffer B, as appropriate) and incubated at room temperature with gentle agitation
10 for 30 minutes. The sample was centrifuged at 700g for 5 minutes at 4°C. The resin was washed twice with 10ml buffer A or B for 10 minutes, resuspended in 1ml buffer A or B and loaded on a disposable column. The resin was washed at either (i) 4°C with 2ml cold buffer A or (ii) room temperature with 2ml buffer B, until the flow-through reached OD₂₈₀ of 0.02-0.06.

The resin was washed with either (i) 2ml cold 20mM imidazole buffer (300mM NaCl, 50mM
15 phosphate buffer, 20mM imidazole, pH 8) or (ii) buffer D (urea 8M, 10mM Tris-HCl, 100mM phosphate buffer, pH 6.3) until the flow-through reached the O.D₂₈₀ of 0.02-0.06. The His-fusion protein was eluted by addition of 700µl of either (i) cold elution buffer A (300mM NaCl, 50mM phosphate buffer, 250mM imidazole, pH 8) or (ii) elution buffer B (urea 8M, 10mM Tris-HCl, 100mM phosphate buffer, pH 4.5) and fractions collected until the O.D₂₈₀ was 0.1. 21µl of each
20 fraction were loaded on a 12% SDS gel.

K) His-fusion proteins renaturation

10% glycerol was added to the denatured proteins. The proteins were then diluted to 20µg/ml using dialysis buffer I (10% glycerol, 0.5M arginine, 50mM phosphate buffer, 5mM reduced glutathione, 0.5mM oxidised glutathione, 2M urea, pH 8.8) and dialysed against the same buffer at 4°C for 12-
25 14 hours. The protein was further dialysed against dialysis buffer II (10% glycerol, 0.5M arginine, 50mM phosphate buffer, 5mM reduced glutathione, 0.5mM oxidised glutathione, pH 8.8) for 12-14 hours at 4°C. Protein concentration was evaluated using the formula:

$$\text{Protein (mg/ml)} = (1.55 \times \text{OD}_{280}) - (0.76 \times \text{OD}_{260})$$

L) His-fusion large-scale purification (ORFs 111-129)

500ml of bacterial cultures were induced and the fusion proteins were obtained soluble in buffer M1, M2 or M3 using the procedure described above. The crude extract of the bacteria was loaded onto a Ni-NTA superflow column (Quiagen) equilibrated with buffer M1, M2 or M3 depending on the solubilization buffer of the fusion proteins. Unbound material was eluted by washing the column with the same buffer. The specific protein was eluted with the corresponding buffer containing 500mM imidazole and dialysed against the corresponding buffer without imidazole. After each run the columns were sanitized by washing with at least two column volumes of 0.5 M sodium hydroxide and reequilibrated before the next use.

10 M) Mice immunisations

20µg of each purified protein were used to immunise mice intraperitoneally. In the case of ORFs 2, 4, 15, 22, 27, 28, 37, 76, 89 and 97, Balb-C mice were immunised with Al(OH)₃ as adjuvant on days 1, 21 and 42, and immune response was monitored in samples taken on day 56. For ORFs 44, 106 and 132, CD1 mice were immunised using the same protocol. For ORFs 25 and 40, CD1 mice were immunised using Freund's adjuvant, rather than Al(OH)₃, and the same immunisation protocol was used, except that the immune response was measured on day 42, rather than 56. Similarly, for ORFs 23, 32, 38 and 79, CD1 mice were immunised with Freund's adjuvant, but the immune response was measured on day 49.

N) ELISA assay (sera analysis)

20 The acapsulated MenB M7 strain was plated on chocolate agar plates and incubated overnight at 37°C. Bacterial colonies were collected from the agar plates using a sterile dracon swab and inoculated into 7ml of Mueller-Hinton Broth (Difco) containing 0.25% Glucose. Bacterial growth was monitored every 30 minutes by following OD₆₂₀. The bacteria were let to grow until the OD reached the value of 0.3-0.4. The culture was centrifuged for 10 minutes at 10000rpm. The supernatant was discarded and bacteria were washed once with PBS, resuspended in PBS containing 0.025% formaldehyde, and incubated for 2 hours at room temperature and then overnight at 4°C with stirring. 100µl bacterial cells were added to each well of a 96 well Greiner plate and incubated overnight at 4°C. The wells were then washed three times with PBT washing buffer (0.1% Tween-20 in PBS). 200µl of saturation buffer (2.7% Polyvinylpyrrolidone 10 in water) was added to each well and the plates incubated for 2 hours at 37°C. Wells were washed

three times with PBT. 200µl of diluted sera (Dilution buffer: 1% BSA, 0.1% Tween-20, 0.1% NaN₃ in PBS) were added to each well and the plates incubated for 90 minutes at 37°C. Wells were washed three times with PBT. 100µl of HRP-conjugated rabbit anti-mouse (Dako) serum diluted 1:2000 in dilution buffer were added to each well and the plates were incubated for 90 minutes at 37°C. Wells were washed three times with PBT buffer. 100µl of substrate buffer for HRP (25ml of citrate buffer pH5, 10mg of O-phenildiamine and 10µl of H₂O) were added to each well and the plates were left at room temperature for 20 minutes. 100µl H₂SO₄ was added to each well and OD₄₉₀ was followed. The ELISA was considered positive when OD₄₉₀ was 2.5 times the respective pre-immune sera.

10 **O) FACScan bacteria Binding Assay procedure.**

The acapsulated MenB M7 strain was plated on chocolate agar plates and incubated overnight at 37°C. Bacterial colonies were collected from the agar plates using a sterile dracon swab and inoculated into 4 tubes containing 8ml each Mueller-Hinton Broth (Difco) containing 0.25% glucose. Bacterial growth was monitored every 30 minutes by following OD₆₂₀. The bacteria were let to grow until the OD reached the value of 0.35-0.5. The culture was centrifuged for 10 minutes at 4000rpm. The supernatant was discarded and the pellet was resuspended in blocking buffer (1% BSA, 0.4% NaN₃) and centrifuged for 5 minutes at 4000rpm. Cells were resuspended in blocking buffer to reach OD₆₂₀ of 0.07. 100µl bacterial cells were added to each well of a Costar 96 well plate. 100µl of diluted (1:200) sera (in blocking buffer) were added to each well and plates incubated for 2 hours at 4°C. Cells were centrifuged for 5 minutes at 4000rpm, the supernatant aspirated and cells washed by addition of 200µl/well of blocking buffer in each well. 100µl of R-Phicoerytrin conjugated F(ab)₂ goat anti-mouse, diluted 1:100, was added to each well and plates incubated for 1 hour at 4°C. Cells were spun down by centrifugation at 4000rpm for 5 minutes and washed by addition of 200µl/well of blocking buffer. The supernatant was aspirated and cells resuspended in 200µl/well of PBS, 0.25% formaldehyde. Samples were transferred to FACScan tubes and read. The condition for FACScan setting were: FL1 on, FL2 and FL3 off; FSC-H threshold:92; FSC PMT Voltage: E 02; SSC PMT: 474; Amp. Gains 7.1; FL-2 PMT: 539; compensation values: 0.

P) OMV preparations

Bacteria were grown overnight on 5 GC plates, harvested with a loop and resuspended in 10 ml 20mM Tris-HCl. Heat inactivation was performed at 56°C for 30 minutes and the bacteria disrupted by sonication for 10 minutes on ice (50% duty cycle, 50% output). Unbroken cells were removed by centrifugation at 5000g for 10 minutes and the total cell envelope fraction recovered by centrifugation at 50000g for 75 minutes. To extract cytoplasmic membrane proteins from the crude outer membranes, the whole fraction was resuspended in 2% sarkosyl (Sigma) and incubated at room temperature for 20 minutes. The suspension was centrifuged at 10000g for 10 minutes to remove aggregates, and the supernatant further ultracentrifuged at 50000g for 75 minutes to pellet the outer membranes. The outer membranes were resuspended in 10mM Tris-HCl, pH8 and the protein concentration measured by the Bio-Rad Protein assay, using BSA as a standard.

Q) Whole Extracts preparation

Bacteria were grown overnight on a GC plate, harvested with a loop and resuspended in 1ml of 20mM Tris-HCl. Heat inactivation was performed at 56°C for 30 minutes.

R) Western blotting

Purified proteins (500ng/lane), outer membrane vesicles (5µg) and total cell extracts (25µg) derived from MenB strain 2996 were loaded on 15% SDS-PAGE and transferred to a nitrocellulose membrane. The transfer was performed for 2 hours at 150mA at 4°C, in transferring buffer (0.3 % Tris base, 1.44 % glycine, 20% methanol). The membrane was saturated by overnight incubation at 4°C in saturation buffer (10% skimmed milk, 0.1% Triton X100 in PBS). The membrane was washed twice with washing buffer (3% skimmed milk, 0.1% Triton X100 in PBS) and incubated for 2 hours at 37°C with mice sera diluted 1:200 in washing buffer. The membrane was washed twice and incubated for 90 minutes with a 1:2000 dilution of horseradish peroxidase labelled anti-mouse Ig. The membrane was washed twice with 0.1% Triton X100 in PBS and developed with the Opti-4CN Substrate Kit (Bio-Rad). The reaction was stopped by adding water.

S) Bactericidal assay

MC58 strain was grown overnight at 37°C on chocolate agar plates. 5-7 colonies were collected and used to inoculate 7ml Mueller-Hinton broth. The suspension was incubated at 37°C on a nutator and let to grow until OD₆₂₀ was 0.5-0.8. The culture was aliquoted into sterile 1.5ml Eppendorf

tubes and centrifuged for 20 minutes at maximum speed in a microfuge. The pellet was washed once in Gey's buffer (Gibco) and resuspended in the same buffer to an OD₆₂₀ of 0.5, diluted 1:20000 in Gey's buffer and stored at 25°C.

50µl of Gey's buffer/1% BSA was added to each well of a 96-well tissue culture plate. 25µl of diluted mice sera (1:100 in Gey's buffer/0.2% BSA) were added to each well and the plate incubated at 4°C. 25µl of the previously described bacterial suspension were added to each well. 25µl of either heat-inactivated (56°C waterbath for 30 minutes) or normal baby rabbit complement were added to each well. Immediately after the addition of the baby rabbit complement, 22µl of each sample/well were plated on Mueller-Hinton agar plates (time 0). The 96-well plate was incubated for 1 hour at 37°C with rotation and then 22µl of each sample/well were plated on Mueller-Hinton agar plates (time 1). After overnight incubation the colonies corresponding to time 0 and time 1 hour were counted.

Table II (page 493) gives a summary of the cloning, expression and purification results.

Example 1

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 1>:

```

1  ATGAAACAGA CAGTCAA.AT GCTTGCCGCC GCCCTGATTG CCTTGGGCTT
51  GAACCGACCG GTGTGGNCGG ATGACGTATC GGATTTTCGG GAAAACTTGC
101 A.GCGGCAGC ACAGGGAAAT GCAGCAGCCC AATACAATTT GGGCGCAATG
151 TAT.TACAAA GGACGCGCGT GCGCCGGGAT GATGCTGAAG CGGTCAGATG
201 GTATCGGCAG CCGGCGGAAC AGGGGTTAGC CCAAGCCCAA TACAATTTGG
251 GCTGGATGTA TGCCAACGGG CGCGC.GTGC GCCAAGATGA TACCGAAGCG
301 GTCAGATGGT ATCGGCAGGC GGCAGCGCAG GGGGTTGTCC AAGCCCAATA
351 CAATTTGGGC GTGATATATG CCGAAGGACG TGGAGTGCGC CAAGACGATG
401 TCGAAGCGGT CAGATGGTTT CGGCAGGCGG CAGCGCAGGG GGTAGCCCAA
451 GCCCAAAACA ATTTGGGCGT GATGTATGCC GAAAGANCGC GCGTGCGCCA
501 AGACCG...
```

This corresponds to the amino acid sequence <SEQ ID 2; ORF37>:

```

1  MKQTVXMLAA ALIALGLNRP VWXDDVSDFR ENLXAAQGN AAAQYNLGAM
51  YXQRTVRVRD DAEAVRWYRQ PAEQGLAQQA YNLGWMYANG RXVRQDDTEA
101 VRWYRQAAQ GVVQAQYNLG VIYAEGRGVR QDDVEAVRWF RQAAQGVAAQ
151 AQNNLGVMYA ERXRVRQD...
```

Further work revealed the complete nucleotide sequence <SEQ ID 3>:

```

1  ATGAAACAGA CAGTCAAATG GCTTGCCGCC GCCCTGATTG CCTTGGGCTT
51  GAACCGAGCG GTGTGGGCGG ATGACGTATC GGATTTTCGG GAAAACTTGC
101 AGGCGGCAGC ACAGGGAAAT GCAGCAGCCC AATACAATTT GGGCGCAATG
151 TATTACAAAAG GACGCGGCGT GCGCCGGGAT GATGCTGAAG CGGTCAGATG
201 GTATCGGCAG CCGGCGGAAC AGGGGTTAGC CCAAGCCCAA TACAATTTGG
251 GCTGGATGTA TGCCAACGGG CGCGGCGTGC GCCAAGATGA TACCGAAGCG
301 GTCAGATGGT ATCGGCAGGC GGCAGCGCAG GGGGTTGTCC AAGCCCAATA
351 CAATTTGGGC GTGATATATG CCGAAGGACG TGGAGTGCGC CAAGACGATG
401 TCGAAGCGGT CAGATGGTTT CGGCAGGCGG CAGCGCAGGG GGTAGCCCAA
451 GCCCAAAACA ATTTGGGCGT GATGTATGCC GAAAGACGCG GCGTGCGCCA
501 AGACCGCGCC CTTGCACAAG AATGGTTTGG CAAGGCTTGT CAAAACGGAG
551 ACCAAGACGG CTGCGACAAT GACCAACGCC TGAAGGCGGG TTATTGA
```

This corresponds to the amino acid sequence <SEQ ID 4; ORF37-1>:

```

      1  MKQTVKWLAA ALIALGLNRA VWADDVSDFR ENLQAAAQGN AAAQYNLGAM
     51  YYKGRGVRRD DAEAVRWYRQ AAEQGLAQAO YNLGWMYANG RGVRRQDDTEA
    101  VRWYRQAAAQ GVVQAQYNLG VIYAEGRGVR QDDVEAVRWF RQAAAQGVAAQ
    151  AQNNLGVMYA ERRGVRQDRA LAQEWFGKAC QNGDQDGCND DQRLKAGY*

```

Further work identified the corresponding gene in strain A of *N.meningitidis* <SEQ ID 5>:

```

      1  ATGAAACAGA CAGTCAAATG GCTTGCCGCC GCCCTGATTG CCTTGGGCTT
     51  GAACCAAGCG GTGTGGGCGG ATGACGTATC GGATTTTCGG GAAAACCTGC
    101  AGGCGGCAGC ACAGGGAATG GCAGCAGCCC AAAACAATTT GGGCGTGATG
    151  TATGCCGAAA GACGCGGCGT GCGCCAAGAC CGCGCCCTTG CACAAGAATG
    201  GCTTGGCAAG GCTTGTCAAA ACGGATACCA AGACAGCTGC GACAATGACC
    251  AACGCCTGAA AGCGGTTTAT TGA

```

This encodes a protein having amino acid sequence <SEQ ID 6; ORF37a>:

```

      1  MKQTVKWLAA ALIALGLNQA VWADDVSDFR ENLQAAAQGN AAAQNNLGVM
     51  YAERRGVRQD RALAQEWLK ACQNGYQDSC DNDQRLKAGY *

```

The originally-identified partial strain B sequence (ORF37) shows 68.0% identity over a 75aa overlap with ORF37a:

```

20      orf37.pep      10      20      30      40      50      60
      MKQTVXMLAAALIALGLNRPVWXDDVSDFRENLXAAAQGNAAAQYNLGAMYXQRTVRVRD
      orf37a          10      20      30      40      50      60
      MKQTVKWLAAALIALGLNQAVWADDVSDFRENLQAAAQGNAAAQNNLGVMYAERRGVRQD

25      orf37.pep      70      80      90      100     110     120
      DAEAVRWYRQPAEQGLAQAYNLGWMYANGRXVRQDDTEAVRWYRQAAAQGVVQAQYNLG
      | | : | : : |
      orf37a          70      80      90
      RALAQEWLKACQNGYQDSCDNDQRLKAGYX

```

30 Further work identified the corresponding gene in *N.gonorrhoeae* <SEQ ID 7>:

```

      1  ATGAAACAGA CAGTCAAATG GCTTGCCGCC GCCCTGATTG CCTTGGGCTT
     51  GAACCAAGCG GTGTGGGCGG GTGACGTATC GGATTTTCGG GAAAACCTGC
    101  AGGCGGCAGC ACAGGGAATG GCAGCAGCCC AATTCAATTT GGGCGTGATG
    151  TATGAAAATG GACAAGGAGT TCGTCAAGAT TATGTACAGG CAGTGCAGTG
    201  GTATCGCAAG GCTTCAGAAC AAGGGGATGC CCAAGCCCAA TACAATTTGG
    251  GCTTGATGTA TTACGATGGA CGCGGCGTGC GCCAAGACCT TCGCTCGCT
    301  CAACAATGGC TTGGCAAGGC TTGTCAAACG GGAGACCAAA ACAGCTGCGA
    351  CAATGACCAA CGCTGAAGG CGGGTTATTA A

```

This encodes a protein having amino acid sequence <SEQ ID 8; ORF37ng>:

```

40      1  MKQTVKWLAA ALIALGLNQA VWAGDVSDFR ENLQAAEQGN AAAQFNLGVM
     51  YENGQGVQRD YVQAVQWYRK ASEQGDAAQAO YNLGLMYIDG RGVRRQDLALA
    101  QQWLKACQON GDQNSCDNDQ RLKAGY*

```

The originally-identified partial strain B sequence (ORF37) shows 64.9% identity over a 111aa overlap with ORF37ng:

```

45      orf37.pep      60
      MKQTVXMLAAALIALGLNRPVWXDDVSDFRENLXAAAQGNAAAQYNLGAMYXQRTVRVRD
      orf37ng          60
      MKQTVKWLAAALIALGLNQAVWAGDVSDFRENLQAAEQGNAAAQFNLGVMYENGQGVQRD

50      orf37.pep      120
      DAEAVRWYRQPAEQGLAQAYNLGWMYANGRXVRQDDTEAVRWYRQAAAQGVVQAQYNLG
      ::||:|:|: :||| ||||| || :|| |||| :| :| :| :|
      orf37ng          120
      YVQAVQWYRKASEQGDAAQAYNLGLMYIDGRGVRRQDLALAQWLKACQNGDQNSCDNDQ

      orf37.pep      168
      VIYAEGRGVRQDDVEAVRWFRQAAAQGVAAQNNLGVMYAERXVRQRD
55      orf37ng          126
      RLKAGY

```

The complete strain B sequence (ORF37-1) and ORF37ng show 51.5% identity in 198 aa overlap:

		10	20	30	40	50	60
	orf37-1.pep	MKQTVKWLAAALIALGLNRAVWADDVSDFRENLQAAQGNAAQYNLGAMYYKGRGVR					
5	orf37ng	MKQTVKWLAAALIALGLNRAVWAGDVSDFRENLQAAEQGNAAQFNLGVMYENGQGV					
		10	20	30	40	50	60
	orf37-1.pep	DAEAVRWYRQAAEQGLAQYNLGWMYANGRGVRQDDTEAVRWYRQAAQGVVQAQYNLG					
10	orf37ng	YVQAVQWYRKASEQGDAAQYNLGLMYDGRGVRQD-----					
		70	80	90			
	orf37-1.pep	VIYAEGRGVRQDDVEAVRWFRQAAQGVQAQNNLGVMYAERRGVRQDRALAQEWFGKAC					
15	orf37ng	-----LALAQQWLKAC					
		130	140	150	160	170	180
						100	
20	orf37-1.pep	QNGDQDQDNDQRLKAGYX					
	orf37ng	QNGDQNSCNDQRLKAGYX					
		110	120				

- 25 Computer analysis of these amino acid sequences indicates a putative leader sequence, and it was predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

- ORF37-1 (11kDa) was cloned in pET and pGex vectors and expressed in *E.coli*, as described above. The products of protein expression and purification were analyzed by SDS-PAGE. Figure 30 1A shows the results of affinity purification of the GST-fusion protein, and Figure 1B shows the results of expression of the His-fusion in *E.coli*. Purified GST-fusion protein was used to immunise mice, whose sera were used for ELISA (positive result), FACS analysis (Figure 1C), and a bactericidal assay (Figure 1D). These experiments confirm that ORF37-1 is a surface-exposed protein, and that it is a useful immunogen.

- 35 Figure 1E shows plots of hydrophilicity, antigenic index, and AMPHI regions for ORF37-1.

Example 2

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 9>:

	TTCCGCGCA	CATCGGCGGT	TTGAAGGTCA	ATGCCCCCGT	CAAATCCGCA
40	GGCGTATTGG	TCGGGCGCGT	CGGCGCTATC	GGACTTGACC	CGAAATCCTA
	TCAGGCGAGG	GTGCGCCTCG	ATTTGACCGG	CAAGTATCAG	TTTCAGCAGCG
	ACGTTTCCGC	GCAAATCCTG	ACTTCsGGAC	TTTGGGCGA	GCAGTACATC
	GGGCTGCAGC	AGGCGGGCGA	CACGGAAAAC	CTTGCTGCCG	GCGACACCAT
	CTCCGTAACC	AGTTCTGCAA	TGGTCTTGGA	AAACCTTATC	GGCAAATTCA
45	TGACGAGTTT	TGCCGAGAAA	AATGCCGACG	GCGGCAATGC	GGAAAAAGCC
	GCCGAATAA				

This corresponds to the amino acid sequence <SEQ ID 10>:

1 FGDIGGLKVN APVKSAGVLV GRVGAIGLDP KSYQARVRLD LDGKYQFSSD
51 VSAQILTSL LGEQYIGLQQ GGDENLAAG DTISVTSSAM VLENLIGKFM

101 TSFAEKNADG GNAEKAAE*

Computer analysis of this amino acid sequence gave the following results:

Homology with a hypothetical *H.influenzae* protein (ybrd.haein; accession number p45029)

SEQ ID 9 and ybrd.haein show 48.4% aa identity in 122 aa overlap:

```

5      20      30      40      50      60      70
ybrd.h LGIGALVFLGLRVANVQGFATKSYTATFDNIGGLKVRAFLKIGGVVIGRVSAITLDE
N.m      FGDIGGLKVNAPVKSAGVLVGRVGAIGLDP
              10      20      30

10     80     90     100     110     120     130
ybrd.h KSYLPKVSIAINQYNEIPENSSLSIKTSGLLGEQYIALTMGFDDGDTAMLNKGSQIQDT
N.m      KSYQARVRLDLGKY-QFSSDVSAQILTSGLLGEQYIGLQQG---GDTENLAAGDTISVT
              40      50      60      70      80

15     140     150     160
ybrd.h TSAMVLEDLIGQFL--YGSKKSDGNEKSESTEQ
N.m      SSAMVLENLIGKFMTSFAEKNADGGNAEKAAEX
              90      100     110     120

```

Homology with a predicted ORF from *N.gonorrhoeae*

SEQ ID 9 shows 99.2% identity over a 118aa overlap with a predicted ORF from *N. gonorrhoeae*:

```

25     20     30     40     50     60     70
ybrd      GAAVAFLAFRVAGGAFFGGSDKTYAVYADFGDIGGLKVNAPVKSAGVLVGRVGAIGLDP
N.m      FGDIGGLKVNAPVKSAGVLVGRVGAIGLDP
              10     20     30

30     80     90     100     110     120     130
ybrd      KSYQARVRLDLGKYQFSSDVSAQILTSGLLGEQYIGLQQGGDTENLAAGDTISVTSSAM
N.m      KSYQARVRLDLGKYQFSSDVSAQILTSGLLGEQYIGLQQGGDTENLAAGDTISVTSSAM
              40     50     60     70     80     90

35     140     150     160
ybrd      VLENLIGKFMTSFAEKNAEAGNAEKAAEX
N.m      VLENLIGKFMTSFAEKNADGGNAEKAAEX
              100     110     120

```

The complete ybrd *H.influenzae* sequence has a leader sequence and it is expected that the full-length homologous *N.meningitidis* protein will also have one. This suggests that it is either a membrane protein, a secreted protein, or a surface protein and that the protein, or one of its epitopes, could be a useful antigen for vaccines or diagnostics.

Example 3

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 11>:

```

50      1      . . ATTTTGATAT ACCTCATCCG CAAGAATCTA GGTTCGCCCC TCTTCTTCTT
      51      TCAGGAACGC CCCGGAAGG ACGGAAAACC TTTTAAATG GTCAAATTC
      101     GTTCCATGCG CGACGGCTTG TATTCAGACG GCATTCCGCT GCCCGACGGA
      151     GAACGCCTGA CACCGTTCGG CAAAAAAGT CGTGCCGCA GTwTGGACGA
      201     ACTGCCTGAA TTATGGAATA TCTTAAAGG CGAGATGAGC CTGGTCGGCC
      251     CCGCCCCGCT GCTGATGCAA TATCTGCCGC GTACGACAA CTCCAAAAC
      301     CGCGCCACG AAATGAAACC CGGCATTACC GGCTGGGCGC AGGTCAACGG

```

5
10

```

351 GCGCAACGCg CTTTCGTGGG ACGAAAAATT CGCCTGCGAT GTTTGGTATA
401 TCGACCACTT CAGCCTGTGC CTCGACATCA AAATCCTACT GCTGACGGTT
451 AAAAAAGTAT TAATCAAGGA AGGGATTTCC GCACAGGGCG AACA.aCCAT
501 GCCCCCTTTC ACAGGAAAAC GCAAACCTCGC CGTCGTCCGT GCGGGCGGAC
551 ACGGAAAAGT CGTTGCCGAC CTTGCCGCCG CACTCGGCCG GTACAGGGAA
601 ATCGTTTTTC TGGACGACCG CGCACAAGGC AGCGTCAACG GCTTTTCCGT
651 CATCGGCACG ACGCTGCTGC TTGAAAACAG TTTATCGCCC GAACAATACG
701 AAGTCGCCGT CGCCGTCCGC AACAAACGCA TCCGCCGCCA AATCGCCGAA
751 AAAGCCGCCG CGCTCGGCTT CGCCCTGCCG GTACTGGTTC ATCCGAGCGC
801 GACCGTCTCG CCTTCTGCAA CAGTCGGACA AGGCAGCGTC GTTATGGCGA
851 AAGCGGTCG..

```

This corresponds to the amino acid sequence <SEQ ID 12; ORF3>:

15

```

1 ..ILIYLIRKNL GSPVFFFQER PGKDGPFFKM VKFRSMRDGL YSDGIPLPDG
51 ERLTPFGKKL RAASXDELPE LWNILKGEMS LVGPRPLLMQ YLPLYDNFQN
101 RRHEMKPGIT GWAQVNGRNA LSWDEKFACD VYIDHFSLC LDIKILLTV
151 KKVLIKEGIS AQGEXTMPFF TGKRKLAVVG AGGHGKVVD LAAALGRYRE
201 IVFLDDRAQG SVNGFSVIGT TLLLENSLSP EQYDVAVAVG NNRIRRQIAE
251 KAAALGFALP VLVHPDATVS PSATVGQGSV VMAKAV..

```

Further sequence analysis revealed the complete nucleotide sequence <SEQ ID 13>:

20
25
30
35
40
45

```

1 ATGAGTAAAT TCTTCAAACG CCTGTTTGAC ATTGTTGCCT CCGCCTCGGG
51 ACTGATTTC CTCTCGCCAG TATTTTGTAT TTTGATATAC CTCATCCGCA
101 AGAATCTAGG TTCGCCCGTC TTCTTCTTTC AGGAACGCCC CGGAAAGGAC
151 GGAAAACCTT TAAAAATGGT CAAATTCCGT TCCATGCGCG ACGCGCTTGA
201 TTCAGACGGC ATTCCGCTGC CCGACGGAGA ACGCCTGACA CCGTTCGGCA
251 AAAAACTGCG TGCCGCCAGT TTGGACGAAC TGCCCTGAAT ATGGAATATC
301 TTAAGAGCG AGATGAGCCT GGTCCGCCCC CGCCCGCTGC TGATGCAATA
351 TCTGCCGTG TACGACAACT TCCAAAACCG CCGCCACGAA ATGAAACCCG
401 GCATTACCGG CTGGGCGCAG GTCAACGGGC GCAACGCGCT TTCGTGGGAC
451 GAAAAATTCG CCTGCGATGT TTGGTATATC GACCACTTCA GCCTGTGCCT
501 CGACATCAAA ATCCTACTGC TGACGGTTAA AAAAGTATTA ATCAAGGAAG
551 GGATTTCCGC ACAGGGCGAA GCCACCATGC CCCCTTTTAC AGGAAAACGC
601 AAACCTCGCG TCGTCCGTGC GGGCGGACAC GGAAAAGTCG TTGCCGACCT
651 TGCCGCCGCA CTCGCCGGT ACAGGGAAT CGTTTTTCTG GACGACCGCG
701 CACAAGGCAG CGTCAACGGC TTTTCCGTCA TCGGCACGAC GCTGCTGCTT
751 GAAAACAGTT TATCGCCCGA ACAATACGAC GTCGCCGTGC CCGTCGGCAA
801 CAACCGCATC CGCCGCCAAA TCGCCGAAAA AGCCGCCGCG CTCGGCTTCG
851 CCTGCCCGT TCTGGTTCAT CCGGACGCGA CCGTCTCGCC TTCTGCAACA
901 GTCGGACAAG GCAGCGTCGT TATGGCGAAA GCCGTCGTAC AGGCAGGCAG
951 CGTATTGAAA GACGGCGTGA TTGTGAACAC TGCCGCCACC GTCGATCAGC
1001 ACTGCCTGCT TAACGCTTTC GTCCACATCA GCCCAGGCGC GCACCTGTGC
1051 GGCAACACGC ATATCGGCGA AGAAAGCTGG ATAGGCACGG GCGCGTCGAG
1101 CCGCCAGCAG ATCCGTATCG GCAGCCGCGC AACCATTGGA GCGGGCGCAG
1151 TCGTCGTACG CGACGTTTCA GACGGCATGA CCGTCGCGGG CAATCCGCGA
1201 AAGCCGTGCG CGCGCAAAAA CCCCAGAGACC TCGACAGCAT AA

```

45 This corresponds to the amino acid sequence <SEQ ID 14; ORF3-1>:

50

```

1 MSKFFKRLFD IVASASGLIF LSPVFLILY LIRKNLGSFV FFFQERPGKD
51 GKPFKMVKFR SMRDALDS DG IPLPDGERLT PFGKKLRAAS LDELPELWNI
101 LKGEMSLVGP RPLLMQYLPL YDNFQNRHE MKPGITGWAQ VNGRNALSWD
151 EKFACDVWYI DHFSLCLDIK ILLLTVKKVL IKEGISAQGE ATMPFFTGKR
201 KLAVVGAGGH GKVVADLAAA LGRYREIVFL DDRAQGSVNG FSVIGTLLLL
251 ENSLSPEQYD VAVAVGNNRI RRQIAEKAAA LGFALPVLVH PDATVSPSAT
301 VGQGSVVMK AVVQAGSVLK DGVIVNTAAT VDHDCLLNAF VHISPGAHL
351 GNTHIGEE SW IGTGACSRQQ IRIISRATIG AGAVVVRDVS DGMTVAGNFA
401 KPLPRKNPET STA*

```

55 Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF3 shows 93.0% identity over a 286aa overlap with an ORF (ORF3a) from strain A of *N. meningitidis*:

15

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25

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35 The complete length ORF3a nucleotide sequence <SEQ ID 15> is:

40

45

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60

This is predicted to encode a protein having amino acid sequence <SEQ ID 16>:

65

301 VGQGGVMAK AVVQADSVLK DGVIVNTAAT VDHDCLLDAF VHISPGAHL S
 351 GNTRIGEESW IGTGACSRQQ IRIGSRATIG AGAVVVRDVS DGMTVAGNPA
 401 KPLAGKNTET LRS*

Two transmembrane domains are underlined.

5 ORF3-1 shows 94.6% identity in 410 aa overlap with ORF3a:

		10	20	30	40	50	60
	orf3a.pep	MSKFFKRLFDIVASASGLIFLSPVFLILYLRKNLGSPVFFFQERPGKDGPVKF					
	orf3-1	MSKFFKRLFDIVASASGLIFLSPVFLILYLRKNLGSPVFFFQERPGKDGPVKF					
10		10	20	30	40	50	60
	orf3a.pep	SMHDALDSDGILLPDGERLTPFGKKLRAASLDELPELWNVLKGDMSLVGPRPLLMQYLPL					
	orf3-1	SMRDALDSDGIPLPDGERLTPFGKKLRAASLDELPELWNILKGEMSLVGPRPLLMQYLPL					
15		70	80	90	100	110	120
	orf3a.pep	SMHDALDSDGILLPDGERLTPFGKKLRAASLDELPELWNVLKGDMSLVGPRPLLMQYLPL					
	orf3-1	SMRDALDSDGIPLPDGERLTPFGKKLRAASLDELPELWNILKGEMSLVGPRPLLMQYLPL					
		70	80	90	100	110	120
	orf3a.pep	YDNFQNRHEMKPGITGWAQVNGRNLSDWERFACDIWYIDHFSCLCDIKILLTVKKVL					
	orf3-1	YDNFQNRHEMKPGITGWAQVNGRNLSDWEKFACDVWYIDHFSCLCDIKILLTVKKVL					
20		130	140	150	160	170	180
	orf3a.pep	YDNFQNRHEMKPGITGWAQVNGRNLSDWERFACDIWYIDHFSCLCDIKILLTVKKVL					
	orf3-1	YDNFQNRHEMKPGITGWAQVNGRNLSDWEKFACDVWYIDHFSCLCDIKILLTVKKVL					
		130	140	150	160	170	180
	orf3a.pep	IKEGISAQGEATMPFFTGKRKLAVVGAGGHGKVVAAELAAALGTYEIVFLDDRVSQSVNG					
	orf3-1	IKEGISAQGEATMPFFTGKRKLAVVGAGGHGKVVADLAAALGRYREIVFLDDRVSQSVNG					
25		190	200	210	220	230	240
	orf3a.pep	IKEGISAQGEATMPFFTGKRKLAVVGAGGHGKVVAAELAAALGTYEIVFLDDRVSQSVNG					
	orf3-1	IKEGISAQGEATMPFFTGKRKLAVVGAGGHGKVVADLAAALGRYREIVFLDDRVSQSVNG					
		190	200	210	220	230	240
	orf3a.pep	FPVIGTTLLENLSLSPQYDVAVAVGNNRIRQAIEKAAALGFALPVLVHPDSTVSFSA					
	orf3-1	FSVIGTTLLENLSLSPQYDVAVAVGNNRIRQAIEKAAALGFALPVLVHPDSTVSFSA					
30		250	260	270	280	290	300
	orf3a.pep	FPVIGTTLLENLSLSPQYDVAVAVGNNRIRQAIEKAAALGFALPVLVHPDSTVSFSA					
	orf3-1	FSVIGTTLLENLSLSPQYDVAVAVGNNRIRQAIEKAAALGFALPVLVHPDSTVSFSA					
		250	260	270	280	290	300
	orf3a.pep	VGQGGVMAKAVVQADSVLKDGVIVNTAATVDHDCLLDAFVHISPGAHLGNTRIGEESW					
	orf3-1	VGQGSVMAKAVVQAGSVLKDGVIVNTAATVDHDCLLNAFVHISPGAHLGNTHIGEESW					
35		310	320	330	340	350	360
	orf3a.pep	VGQGGVMAKAVVQADSVLKDGVIVNTAATVDHDCLLDAFVHISPGAHLGNTRIGEESW					
	orf3-1	VGQGSVMAKAVVQAGSVLKDGVIVNTAATVDHDCLLNAFVHISPGAHLGNTHIGEESW					
40		310	320	330	340	350	360
	orf3a.pep	VGQGGVMAKAVVQADSVLKDGVIVNTAATVDHDCLLDAFVHISPGAHLGNTRIGEESW					
	orf3-1	VGQGSVMAKAVVQAGSVLKDGVIVNTAATVDHDCLLNAFVHISPGAHLGNTHIGEESW					
		310	320	330	340	350	360
	orf3a.pep	IGTGACSRQQIRIGSRATIGAGAVVVRDVS DGMTVAGNPAKPLAGKNTETLRSX					
	orf3-1	IGTGACSRQQIRIGSRATIGAGAVVVRDVS DGMTVAGNPAKPLPRKNPETSTAX					
45		370	380	390	400	410	
	orf3a.pep	IGTGACSRQQIRIGSRATIGAGAVVVRDVS DGMTVAGNPAKPLAGKNTETLRSX					
	orf3-1	IGTGACSRQQIRIGSRATIGAGAVVVRDVS DGMTVAGNPAKPLPRKNPETSTAX					
		370	380	390	400	410	

Homology with hypothetical protein encoded by *yvfc* gene (accession Z71928) of *B. subtilis*

ORF3 and YVFC proteins show 55% aa identity in 170 aa overlap (BLASTp):

50	ORF3	3	IYLIRKNLGSPVFFFQERPGKDGPVKFMSMRDGLYSDGIPLPDGERLTPFGKKLRA	62
	yvfc	27	I VVRLKIGSPVFFKQVRPGLHGKPF TLYKFR TMTDERDSKGNLLPDEVRLTKTGRLIRK	86
			I ++R +GSPVEF Q RPG GKPF + KFR+M D S G LPD RLT G+ +R	
55	ORF3	63	ASXDELPELWNILKGEMSLVGPRPLLMQYLPLYDNFQNRHEMKPGITGWAQVNGRNL	122
	yvfc	87	LSIDELPQLLVNLKGDLSLVGPRPLMDYLPYTEKQARRHEVKPGITGWAQINGRNAIS	146
			S DELP+L N+LKG++SLVGPRPLM YLPLY Q RRHE+KPGITGWAQ+NGRNA+S	
	ORF3	123	WDEKFACDVWYIDHFSCLDXXXXXXXXXXXXXXXXXEGISAQGEEXTMPFFT	172
	yvfc	147	WEKKFELDVWYVDNWSFFLDLKLCLTVRKVLVSEGIQQTNHVTAERFTG	196
			W++KF DVWY+D++S LD EGI T FTG	

Homology with a predicted ORF from *N.gonorrhoeae*

ORF3 shows 86.3% identity over a 286aa overlap with a predicted ORF (ORF3.ng) from *N. gonorrhoeae*:

5	orf3	ILIIYLIRKNLGSPVFFQERPGKDGKPFKMVKFR	34
	orf3ng	MSKAVKRLFDIIASASGLIVLSPVFLVLIYLIRKNKGSPVFFIRERPGKDGKPFKMVKFR	60
10	orf3	SMRDGLYSDGIPLDPGERLTPFGKKLRAASXDELPELWNILKGEMSLVGPRLPLMQYLPL	94
	orf3ng	SMRDALDSGDIPLPDSERLTDGFKKLRLATSLDELPELWNVLKGEMSLVGPRLPLMQYLPL	120
15	orf3	YDNFQNRHMKPGITGWAQVNGRNLASWDEKFCADVWYIDHFSCLDIKILLTLVKKVL	154
	orf3ng	YNKFQNRHMKPGITGWAQVNGRNLASWDEKFCADVWYIDHFSCLDIKILLTLVKKVL	180
20	orf3	IKEGISAQGEEXTMPFFTGRKRLAVVGAGGHGKVVADLAAALGRYREIVFLDDRAQGSVNG	214
	orf3ng	IKEGISAQGEATMPFFAGNRKLAVIGAGGHGKVVAEALAAALGTIGEIVFLDDRTQGSVNG	240
25	orf3	FSVIGTTLLENLSLPEQYDVAVAVGNRRIRRQIAEKAAALGFALPVLVHPDATVSPSAT	274
	orf3ng	FPVIGTTLLENLSLPEQFDITVAVGNRRIRRQITENAAALGFALPVLVHPDATVSPSAI	300
25	orf3	VGQGSVVMKAV	286
	orf3ng	IGQGSVVMKAVVQAGSVLKDGIVNTAATVDHDCLLDAFVHISPGAHLSGNTRIGEEER	360

The complete length ORF3ng nucleotide sequence <SEQ ID 17> is:

1	ATGAGTAAAG	CCGTCAAACG	CCTGTTGCGAC	ATCATCGCAT	CCGCATCGGG
51	GCTGATTGTC	CTGTCGCCCG	TGTTTTTGGT	TTTAATATAC	CTCATCCGCA
101	AAAACCTTAGG	TTCGCCCGTC	TTCTTCattC	GGGAACGCCc	cgGAAAGGAc
151	ggaaaaacCTT	TTAAATGGT	CAAATTCGGT	TCCAtgcgcg	acgcgcttGA
201	TTCAGACGGC	ATTCCGCTGC	CCGATAGCGA	ACGCCTGACC	GATTTCCGGCA
251	AAAAATTACG	CGCCACCACT	TTGGACGAAC	TTCTTGAATT	ATGGAATGTC
301	CTCAAAGGCG	AGATGAGCCT	GGTCGGCCCC	CGCCCGCTTT	TGATGCAGTA
351	TCTGCCGCTT	TACAACAAT	TTCAAACCG	CGCCACGAA	ATGAAACCGG
401	GCATTACCGG	CTGGGCGCAG	GTCAACGGGC	GCAACGCGCT	TTCTGGGAC
451	GAAAAGTTCT	CCTGCGATGT	TTGGTACACC	GACAATTTCA	GCTTTTGGCT
501	GGATATGAAA	ATCCTGTTTC	TGACAGTCAA	AAAAGTCTTG	ATTAAAGAAG
551	GCATTTCGGC	GCAAGGGGAA	GCCACCATGC	CCCCTTTCGC	GGGGAATCGC
601	AAACTCGCCG	TTATCGGCGC	GGGCGGACAC	GGCAAAGTCG	TTGCCGAGCT
651	TGCCGCGGCA	CTCGGCACAT	ACGGCGAAAT	CGTTTTTCTG	GACGACCGCA
701	CCCAAGCGAG	CGTCAACGGC	TTCCCGCTCA	TCGGCACGAC	GCTGCTGCTT
751	GAAAACAGTT	TATCGCCCGA	ACAATTCGAC	ATCACCGTCG	CCGTCGGCAA
801	CAACCGCATC	CGCCGCCAAA	TCACCGAAAA	CGCCGCGCGC	CTCGGCTTCA
851	AACTGCCCGT	TCTGATTTCAT	CCCGACGCGA	CCGTCTCGCC	TTCTGCAATA
901	ATCGGACAAG	GCAGCGTCGT	AATGGCGAAA	GCCGTCGTAC	AGGCCGGCAG
951	CGTATTGAAA	GACGGCGTGA	TTGTGAACAC	TGCCGCCACC	GTCGATCACG
1001	ACTGCCTGCT	TGACGCTTTC	GtccaCATCA	GCCCGGGCGC	GCACCTGTCTG
1051	GGCAACACGC	GTATCGGCGA	AGAAAGCCGG	ATAGGCACGG	GCGCGTGCAG
1101	CGGCCAGCAG	ACAACCGTCG	GCAGCGGGGT	TACCGccgGT	GCAGGGgcGG
1151	TTATCGTATG	CGACATCCCG	GACGGCATGA	CCGTCTCGGG	CAACCCGGCA
1201	AAGCCCCTTA	CGGGCAAAAA	CCCCAAGACC	GGGACGGCAT	AA

This encodes a protein having amino acid sequence <SEQ ID 18>:

1	MSKAVKRLFD	IIASASGLIV	LSPVFLVLIY	LIRKNLGSPV	FFIRERPGKD
51	GKPFKMVKFR	SMRDALDSDG	IPLPDSERLT	DFGKKLRATS	LDELPELWNV
101	LKGEMSLVGP	RPLLMQYLPL	YNKFQNRHE	MKPGITGWAQ	VNGRNLASWD
151	EKFSCDVWYT	DNFSFWLDMK	ILFLTVMKVL	IKEGISAQGE	ATMPFFAGNR
201	KLAVIGAGGH	GKVVAEALAA	LGTYGEIVFL	DDRTQGSVNG	FPVIGTLLLL
251	ENSLSPEQFD	ITVAVGNRI	RRQITENAAA	LGFKLPVLIH	PDATVSPSAI
301	IGQGSVVMKAV	AVVQAGSVLK	DGVIVNTAAT	VDHDCLLDAF	VHISPGAHL
351	GNTRIGEEER	IGTGACSRQQ	TTVGSGVTAG	AGAVIVCDIP	DGMTVAGNPA
401	KPLTGKNPKT	GTA*			

This protein shows 86.9% identity in 413 aa overlap with ORF3-1:

5	orf3-1.pep	MSKFFKRLFDIVASASGLIFLSPVFLILYLRKNLGSPVFFQERPGKDGPFKMVKFR	10	20	30	40	50	60
	orf3ng	MSKAVKRLFDIIASASGLIVLSPVFLVLIYLRKNLGSPVFFIRERPGKDGPFKMVKFR	10	20	30	40	50	60
10	orf3-1.pep	SMRDALDSGDIPLPDGERLTPFGKKLRAASLDELPELWNILKGEMSLVGPRPLLMQYLPL	70	80	90	100	110	120
	orf3ng	SMRDALDSGDIPLPDSERLTDFGKKLRATSLDELPELWNVLKGEMSLVGPRPLLMQYLPL	70	80	90	100	110	120
15	orf3-1.pep	YDNFQNRHHEMKPGITGWAQVNGRNALSWEKFCADVWYIDHFSCLCDIKILLTVKKVL	130	140	150	160	170	180
	orf3ng	YNKFQNRHHEMKPGITGWAQVNGRNALSWEKFCADVWYIDHFSCLCDIKILLTVKKVL	130	140	150	160	170	180
20	orf3-1.pep	IKEGISAQGEATMPPFTGKRKLAVVGAGGHGKVVADLAAALGRYREIVFLDDRAQGSVNG	190	200	210	220	230	240
	orf3ng	IKEGISAQGEATMPPFAGNRKLAVIGAGGHGKVVAEALAAALGTYGEIVFLDRTQGSVNG	190	200	210	220	230	240
25	orf3-1.pep	FSVIGTLLLENSLSPEQYDVAVAVGNNRIRQIAEKAAALGFALPVLVHPDATVSPSAT	250	260	270	280	290	300
	orf3ng	FPVIGTLLLENSLSPEQFDITVAVGNNRIRQITENAAALGFKLPVLIHPDATVSPSAI	250	260	270	280	290	300
35	orf3-1.pep	VGQGSVMAKAVVQAGSVLKDGVIVNTAATVDHDCLLNAFVHISPGAHLNTHIGEEESW	310	320	330	340	350	360
	orf3ng	IGQGSVMAKAVVQAGSVLKDGVIVNTAATVDHDCLLDAFVHISPGAHLNTRIGEEESR	310	320	330	340	350	360
40	orf3-1.pep	IGTGACSRQQIRIGSRATIGAGAVVVRDVS DGMTVAGNPAKPLPRKNPETSTAX	370	380	390	400	410	
	orf3ng	IGTGACSRQQTTVGSGVTAGAGAVIVCDIPDGMTVAGNPAKPLTGKNPKTGATX	370	380	390	400	410	

In addition, ORF3ng shows significant homology with a hypothetical protein from *B.subtilis*:

45	gnl PID e238668 (Z71928) hypothetical protein [Bacillus subtilis]	
	>gi 1945702 gnl PID e313004 (Z94043) hypothetical protein [Bacillus subtilis]	
50	>gi 2635938 gnl PID e1186113 (Z99121) similar to capsular polysaccharide biosynthesis [Bacillus subtilis] Length = 202	
	Score = 235 bits (594), Expect = 3e-61	
55	Identities = 114/195 (58%), Positives = 142/195 (72%)	
	Query: 5 VKRLFDIIASASGLIVLSPVFLVLIYLRKNLGSPVFFIRERPGKDGPFKMVKFRSMRD 64	
60	+KRLFD+ A+ L S + L I ++R +GSPVFF + RPG GKPF + KFR+M D	
	Sbjct: 3 LKRLFDLTAAIFLLCCTSVIILEFTIAVRLKIGSPVFFKQVRPGLHGKPFETLYKFRTMTD 62	
65	Query: 65 ALDSGDIPLPDSERLTDFGKKLRATSLDELPELWNVLKGEMSLVGPRPLLMQYLPLYNKF 124	
	DS G LPD RLT G+ +R S+DELP+L NVLKG++SLVGPRPLLM YLPLY +	
70	Sbjct: 63 ERDSKGNLLPDEVRLTKTGRILRKLSDIDLPQLNVLKGDLSLVGPRPLLMQYLPLYTEK 122	
75	Query: 125 QNRRHEMKPGITGWAQVNGRNALSWEKFCADVWYIDHFSCLCDIKILLTVKKVLKEG 184	
	Q RRHE+KPGITGWAQ+NGRNA+SW++KF DWY DN+SF+LD+KIL LTV+KVL+ EG	
80	Sbjct: 123 QARRHEVKPGITGWAQINGRNAISWEKKFELDQVYVDNWSFFLDLKLCLTVRKVLVSEG 182	
85	Query: 185 ISAQGEATMPPFAGN 199	
	I T F G+	
90	Sbjct: 183 IQQTNHVTAERFTGS 197	

The hypothetical product of *yyfc* gene shows similarity to EXOY of *R.meliloti*, an exopolysaccharide production protein. Based on this and on the two predicted transmembrane regions in the homologous *N.gonorrhoeae* sequence, it is predicted that these proteins, or their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

5 Example 4

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 19>:

```

1  ..AACCATATGG CGATTGTCAT CGACGAATAC GCGGGCACAT CCGGCTTGGT
51  CACCTTTGAA GACATCATCG AGCAAATCGT CGGCGAAATC GAAGACGAGT
101 TTGACGAAGA CGATAGCGCC GACAATATCC ATGCCGTTTC TTCAGACACG
151 TGGCGCATCC ATGCAGCTAC CGAAATCGAA GACATCAACA CCTTCTTCGG
201 CACGGAATAC AGCATCGAAG AAGCCGACAC CATT.GGCGG CCTGGTCATT
251 CAAGAGTTGG GACATCTGCC CGTGGCGCGG GAAAAAGTCC TTATCGGCGG
301 TTTGCAGTTC ACCGTCGCAC GCGCCGACAA CCGCCGCCTG CATACGCTGA
351 TGGCGACCCG CGTGAAGTAA GC..... ACCGC CGTTTCTGCA
15 401 CAGTTTAG

```

This corresponds to amino acid sequence <SEQ ID 20; ORF5>:

```

1  ..NHMAIVIDEY GGTSGLVTFE DIIEQIVGEI EDEFDEDDSA DNIHAVSSDT
51  WRIHAATEIE DINTFFGTEY SIEEADTI XR PGHSRVGTSA RARRKSPYRR
101 FAVHRRTRRQ PPPAYADGDP REVS....XR RFCTV*

```

20 Further sequence analysis revealed the complete DNA sequence to be <SEQ ID 21>:

```

1  ATGGACGGCG CACAACCGAA AACGAATTTT TTTGAACGCC TGATTGCCCG
51  ACTCGCCCGC GAACCCGATT CCGCCGAAGA CGTATTAAAC CTGCTTCGGC
101 AGGCGCACGA GCAGGAAGTT TTTGATGCGG ATACGCTTTT AAGATTGGAA
151 AAAGTCCTCG ATTTTTCGA TTTGGAAGTG CGCGACGCGA TGATTACGCG
25 201 CAGCCGTATG AACGTTTAA AAGAAAACGA CAGCATCGAG CGCATCACCG
251 CCTACGTTAT CGATACCGCC CATTGCGGCT TCCCCGTCAT CGGCGAAGAC
301 AAAGACGAAG TTTTGGGCAT TTTGCACGCC AAAGACCTGC TCAAAATATAT
351 GTTTAACCCC GAGCAGTTCC ACCTCAAATC CATTCTCCGC CCCGCCGTCT
401 TCGTCCCGCA AGGCAAATCG CTGACCGCCC TTTTAAAAGA GTTCCGCGAA
30 451 CAGCGCAACC ATATGGCGAT TGTCATCGAC GAATACGGCG GCACATCCGG
501 CTTGGTCACC TTTGAAGACA TCATCGAGCA AATCGTCGGC GAAATCGAAG
551 ACGAGTTTGA CGAAGACGAT AGCGCCGACA ATATCCATGC CGTTTCTTCC
601 GAACGCTGGC GCATCCATGC AGCTACCGAA ATCGAAGACA TCAACACCTT
651 CTTGCGCACG GAATACAGCA GCGAAGAAGC CGACACCATT CGGCCTGGTC
35 701 ATTCAGAGAT TGGGACATCT GCCCGTGCGC GGCGAAAAAG TCCTTATCGG
751 CGGTTTGAG TTCACGTCG CACGCGCCGA CAACCGCCGC CTGCATACGC
801 TGATGGCGAC CCGCGTGAAG TAAGCACCGC CGTTTCTGCA CAGTTTAGGA
851 TGACGTACG GCGTTTTTCT GTTTCAATCC GCCCATCCG CCAAACATAA

```

This corresponds to amino acid sequence <SEQ ID 22; ORF5-1>:

```

40 1  MDGAQPKTNF FERLIARLAR EPDSAEDVLN LLRQAHEQEV FDADTLRLLE
51  KVLDFSDLEV RDAMITRSRM NVLKENDSIE RITAYVIDTA HSRFPVIGED
101 KDEVLGILHA KDLLKYMENP EQFHLKSILR PAVFVPEGKS LTALLKEFRE
151 QRNHMAIVID EYGGTSGLVF FEDIIEQIVG EIEDEFDEDD SADNIHAVSS
201 ERWRIHAATE IEDINTFFGT EYSSEEADTI RPHGSRVGTS ARARRKSPYR
45 251 REAVHRRTRR QPPPAYADGD PREVSTAVSA QFRMTVRAFS VSIRPRTQT*

```

Further work identified the corresponding gene in strain A of *N.meningitidis* <SEQ ID 23 >:

```

1  ATGGACGGCG CACAACCGAA AACAAATTTT TTNNAACGCC TGATTGCCCG
51  ACTCGCCCGC GAACCCGATT CCGCCGAAGA CGTATTGACC CTGTTGCGCC
101 AAGCGCACGA ACAGGAAGTA TTTGATGCGG ATACGCTTTT AAGATTGGAA
151 AAAGTCCTCG ATTTTCTGTA TTTGGAAGTG CGCGACGCGA TGATTACGCG
201 CAGCCGTATG AACGTTTAA AAGAAAACGA CAGCATCGAA CGCATCACCG
251 CCTACGTTAT CGATACCGCC CATTGCGGCT TCCCCGTCAT CGGTGAAGAC
301 AAAGACGAAG TTTTGGGTAT TTTGCACGCC AAAGACCTGC TCAAAATATAT
351 GTTCAACCCC GAGCAGTTCC ACCTCAAATC GATATTGCGC CCTGCCGTCT

```

10

401	TCGTCCCCGA	AGGCAAATCG	CTGACCGCCC	TTTTAAAAGA	GTTCCGCGAA
451	CAGCGCAACC	ATATGGCAAT	CGTCATCGAC	GAATACGGCG	GCACGTCGGG
501	TTTGGTAACT	TTTGAAGACA	TCATCGAGCA	AATCGTCGGC	GACATCGAAG
551	ATGAGTTTGA	CGAAGACGAA	AGCGCGGACA	ACATCCACGC	GGTTTTCCGC
601	GAACGCTGGC	GCATCCACGC	GGCTACCGAA	ATCGAAGACA	TCAACGCCCT
651	TTTCGGCACG	GAATACAGCA	GCGAAGAAGC	CGACACCATC	GGCGCGCNTG
701	GTCAATTGAG	AATTGGNACA	CCTGCCCGTG	CGCGCGGAAA	AAGTCNTTAT
751	CGGCGNNTTG	CANTTCAACG	TCGCCNCGCG	NGACAACCGC	CGCCTGCATA
801	CGCTGATGGC	GACCCGCGTG	AAGTAAGCTC	CGCCGTTTCT	GTACAGTTTA
851	GGATGACGGT	ACGGGCGGTT	TCTGTTTCAA	TCCGCCCCAT	CCGCCANACA
901	TAA				

15

```

1 MDGAQPKTNF XXRLIARLAR EPDSAEDVLT LLRQAHEQEV FDADTLLRLR
51 KVLDFSDLEV RDAMITRSM NVLKENDSIE RTIAYIVGTA HSRFPVIGED
101 KDEVLGILHA KDLLKYMFRM EQFHLKSILR PAVFVPEGKS LTALLKEFRE
151 QRNHMAIVID EYGGTSGLVT FEDIEIQIVG DIEDFEDEBE SADNIHVASA
201 ERWRIHAATE IEDINAFFGT EYSSEEADTI GGXGHSIGIT PARARRKSKY
251 RRAXHXRXR XQPPPAYADG DPREVSSAVS VQFRMTVRAF SVSIRPIRXT
301 *

```

20 The originally-identified partial strain B sequence (ORF5) shows 54.7% identity over a 124aa overlap with ORF5a:

```

                                10          20          30
    orf5.pep                     NHMAIVIDEYGGTSGLVTFEDII EQIVGEI
25   orf5a       FHLKSILRPVAVFVPEGKSLTALLKEFREQRNHMAIVIDEYGGTSGLVTFEDII EQIVGD
                        130      140      150      160      170      180

                                40          50          60          70          80          90
    orf5.pep     EDEFDEDDSDADNIHAVSSDTWRIHAATEIEDINTFFGT EYSIEEADTXRPGHSRVGTSA
    orf5a         EDEFDEDESADNIHVAERWR IHAATEIEDINAF GTEYSSEEADTI GGXGHSGIGTPA
                        190      200      210      220      230      240

                                100        110        120        130
35   orf5.pep     RARRKSPYRRFAVHRRT RRQPPPAYADGDPREV SXXXXXR RCTV
    orf5a         RARRKSXYRRAXHXRXRX QPPPAYADGDPREV SSAYSVSQFRMT VRAFSVSIRPI RTX
                        250      260      270      280      290      300

```

The complete strain B sequence (ORF5-1) and ORF5a show 92.7% identity in 300 aa overlap:

40	orf5a.pep	MDGAQPKTNFXXRLIARLAREPDSAEVDVLTLLRQAHEQEVEFDADTLRLREKVLDFSDLEV
	orf5-1	MDGAQPKTNFFERLIARLAREPDSAEVDVNLNLRQAHEQEVEFDADTLRLREKVLDFSDLEV
45	orf5a.pep	RDAMITSRSMNVLKENDSIERITAYVIDTAHSRFPVIGEDKDEVLGTLHAKDLLKYMENF
	orf5-1	RDAMITSRSMNVLKENDSIERITAYVIDTAHSRFPVIGEDKDEVLGILHAKDLLKYMENF
50	orf5a.pep	EQFHLKSILRPAVFPVEGKSLTALLKEFREQRNHMAIVIDEYGGTSGLVTFEDIEIQIVG
	orf5-1	EQFHLKSILRPAVFPVEGKSLTALLKEFREQRNHMAIVIDEYGGTSGLVTFEDIEIQIVG
55	orf5a.pep	DIEDEFDEDESADNIHAVSAERWRIHAATEIEDINAFFGTEYSSEEADTIGGXGHSIGT
	orf5-1	EIEDEFDEDDSDNIHAVSSERWRIHAATEIEDINTFFGTEYSSEEADTIRP-GHSRVGT
60	orf5a.pep	
	orf5-1	

	1	MDGAQPKTNF	FERLIARLAR	EPDSAEDVLN	LLRQAEQEV	FDADTLTRLE
	51	KVLDFAELE	RDAMITRSRM	NVLKENDSIE	RITAYVIDTA	HSRFPVIGED
10	101	KDEVLGLIHA	KDLLKYMFP	EQFHLKSVLR	PAVFVPEGKS	LTALLKEFRE
	151	QRNHMAIVID	EYGGTSGLVT	FEDIIEQIVG	DIEDEFDEDE	SADDIHSVSA
	201	ERWRIHAAE	IEDINAFFGT	EYGSEEDATI	RRRHSG*GT	PARARRKSPY
	251	RRFAVHRRR	ROPPPAHADG	DPREVSRACP	HLRFCTV*	

15	1	ATGGACGGCG	CACAACCGAA	AACAAATTTT	TTTGAACGCC	TGATTGCCCG
	51	ACTCGCCCGC	GAACCCGATT	CGGCCGAAGA	CGTATTAAAC	CTGCTTCGGC
	101	AGGCCGCAGCA	ACAGGAAGTT	TTTGATGCCG	ACACACTGAC	CCGGCTCGGAA
	151	AAAGTATTGG	ACTTTGCCGA	GCTGGAAGTG	CGCGATCGAA	TGATTACGGC
	201	CAGCCGCATG	AACGTATTGA	AAGAAAACGA	CAGCATCGAA	CGCATCACCG
20	251	CCTACGTCAT	CGATACCGCC	CATTTCGCGT	TCCCCGTCAT	CGGCGAAGAC
	301	AAAGACGAAG	TTTTGGGCAT	TTTGCACGCG	AAAGACCTGC	TCAAATATAT
	351	GTTCAACCCC	GAGCAGTTCC	ACCTGAAATC	CGTCTTGGCG	CTTGCGGTTT
	401	TCGTGCCCGA	AGGCAAATCT	TTGACCGCCC	TTTTAAAAGA	GTTCCGCGAA
	451	CAGCGCAACC	ATATGGCAAT	CGTCATCGAC	GAATACGGCG	GACAGTCGGG
25	501	TTTGGTCACC	TTTGAAGACA	TCATCGAGCA	AATCGTCGGT	GCATCTCGAG
	551	ACGAGTTTGA	CGAAGACGAA	AGCGcgcagc	acatCCACTC	cgTTTccgCC
	601	GAACGCTGGC	GCATCCacgc	ggctaCCGAA	ATCGAAGaca	TCAACGCCTT
	651	TTTCGGTACG	GAatacggca	gcgaagaagc	cgacaccatc	cgggcggtTG
	701	GTCATTACG	AATTGGGACA	CCTGCCCGTG	CGCGGCGAAA	AAGTCTTATA
30	751	gggcgGTTTG	Cagttcacgc	tCGCCCGCGC	CGACAACCGC	CGCCTGCACA
	801	CGCTGATGGC	GACCCGCGTG	AAGTAAGCAG	AGCCTGCCcg	AccgcggttT
	851	CTGCacAGTT	TAGGatgACG	gtaCGGTCGT	TTTCTGTTTC	AATCCGCCCC
	901	ATCCGCCAAA	CATAA			

35	1	MDGAQPKTNF	FERLIARLAR	EPDSAEDVLN	LLRQAHEQEV	FDADTLTRLE
	51	KVLDFÆLEV	RDAMITRSM	NVLKENDSIE	RITAYVIDTA	HSRFPVIGED
	101	KDEVLGILHA	KDLLKYMFPN	EQFHLKSQVL	PAVEVPEGKS	LTALLKEFRE
	151	QRNHMAIVID	YGGTSGGLVT	FEDIIEQIVG	DIEDEFEDE	SADDIHSVSA
	201	ERWRIHAATE	IEDINAFFGT	EGYSEEADTI	RRLGHSIGIT	PARARRKSPY
40	251	RRFAVHRPR	RQPPPAHADG	DPREVSRACP	TAVSAQFRMT	VRSFSVSIRP
	301	TROT*				

	orf5	NHMAIVIDEYGGTSGLVTFEDIIEQIVGEI	30
45	orf5ng		
	orf5ng	FHLKSVLRPAVFVPEGKSLTALLKEFREQRNHMAIVIDEYGGTSGLVTFEDIIEQIVGDI	182
	orf5	EDEFDEDDSDADNIHAVSSDTWRIHAATEIEDINTFFGTEYSIEEADTIXRPGHSRVGTS	90
50	orf5ng	: : : : : : :	
	orf5ng	EDEFDEDESADDIHSVAERWRIHAATEIEDINAFFGTEYGGSEEADTIRRLGHSGIGTPA	242
	orf5	RARRKSPYRRFAVHRRTRRQPPAYADGDPREVSX----RRFCTV	131
	orf5ng	: : :	
	orf5ng	RARRKSPYRRFAVHRRPRROPPEAHADGDPREVSRRACPHRRFCTV	287

orf5nq-1.pep MDGAQPKTNFFERLIARLAREPDSAEVDVNLNLLROAHEQEVFDADTLTRLEKVLDFAELEV

35 identified the following homologies:

ORF5 and TlyC proteins show 58% aa identity in 77 aa overlap (BLASTp).

40

ORF5	2	HMAIVIDEYGGTSGLVTFEDIIEQIVGEIEDEFDEDDSDADNIHAVSSDTWRIHAATEIED	61
		HMAIV+DE+G SGLVT EDI+EQIVG+IEDEFDE++ AD I +S T+ + A T+I+D	
TlyC	166	HMAIVVDEFGAVSGLVTTIEDILEQIVGDIEDEFDEEEIAD-IRQLSRHTYAVRALTDIDD	224
ORF5	62	INTFFGTEYSIEEADTI	78
		N F T++ EE DTI	
TlyC	225	FNAQENTDFDDEEVDTI	241

45 ORF5ng-1 also shows significant homology with TlyC:

```

SCORES          Init1:   301 Initn:   419 Opt:    668
Smith-Waterman score: 668;      45.9% identity in 242 aa overlap


50              10           20           30           40           50
orf5ng-1.pep     MDGAQPKTNFFERLIARLAR-EPDSAEDVLNLLRQAHEQEVDADTLTRLEK
                  | ||| :||: : | : |:~::~|:~::~~::| :| :|
tlyc_haein       MNDEQQNSNQSENTKKPFFQSLSFGRFFQGELKNREELVEVIRDSEQNLDLIDQNTREMIEG
                  10           20           30           40           50           60


55              60           70           80           90          100          109
orf5ng-1.pep     VLDFAEAEVRDAMITRSRMNVLKENDSIERTAYVIDTAHSRFPVIGE--DKDEVLGILH
                  ::::::|:|:| || ||: ~::~~:: ~::|:~::~|:~::~~::|:~::~|:~::~|:~::~|:~::~|
tlyc_haein       VMEIAELRVRDIMIPRSQIIFIEDQQDLNTCLNTIIESAHSRFPVIADADDNRDNIVIGILH
                  70           80           90          100          110          120


60              110          120          130          140          150          160
orf5ng-1.pep     AKDLLKYMF-NPEQFHLSVLRPAVFVPEGKSLTALLKEFREQRNHMAIVIDEYGGTSGGL
                  ||||~::~~::|:~::~|:~::~|:~::~|:~::~|:~::~|:~::~|:~::~|:~::~|:~::~|:~::~|:~::~|:~::~|
tlyc_haein       AKDLLKFLREDAEVFDLSSLLRPVVIVPESKRVDRLMKDFRSERFHMAIVVDEFQAVSGGL

```

		130	140	150	160	170	180
	170	180	190	200	210	220	
5	orf5ng-1.pep	VTFEDIIEQIVGDIIEDEFDEDESADDIHSVSAERWRIHAATEIEDINAFFGTEYGSEED					
	tlyc_haein	VTIEDILEQIVGDIIEDEFDEEEIAD-IRQLSRHTYAVRALTDIDDFNAQFNTDFDDEEVD					
		190	200	210	220	230	
10		230	240	250	260	270	280
	orf5ng-1.pep	TIRRLGHSGIG-TPARARRKSPYRRFAVHRRPRRQPPPAHADGDPREVSACPTAVSAQF					
	tlyc_haein	TIGGLIMQTFGYLPKRGEIILKNLQFKVTSADSRRLIQLRVTVPDEHLAEMNNVDEKSE					
		240	250	260	270	280	290

15 Homology with a hypothetical secreted protein from *E.coli*:

ORF5a shows homology to a hypothetical secreted protein from *E.coli*:

20 sp|P77392|YBEX_ECOLI HYPOTHETICAL 33.3 KD PROTEIN IN CUTE-ASNB INTERGENIC REGION
>gi|1778577 (U82598) similar to *H. influenzae* [*Escherichia coli*] >gi|1786879
(AE000170) f292; This 292 aa ORF is 23% identical (9 gaps) to 272 residues of an
approx. 440 aa protein YTFL_HAEIN SW: P44717 [*Escherichia coli*] Length = 292

Score = 212 bits (533), Expect = 3e-54
Identities = 112/230 (48%), Positives = 149/230 (64%), Gaps = 3/230 (1%)

25 Query: 2 DGAQPKTNFXRLIARLAR-EPDSAEDVLTLLRQAHEQEVFDADTLLRLEKVLDFSDLEV 60
D K F L+++L EP + +++L L+R + + ++ D DT LE V+D +D V
Sbjct: 10 DTISNKKGFFSLLLSQLFHGEPKNRDELLALIRDSGQNDLIDEDTRDMLEGVMDIADQRV 69

30 Query: 61 RDAMITRSRMNVLKENDSIERITAYVIDTAHSRFPVIGEDKDEVLGILHAKDLLKYM-FN 119
RD MI RS+M LK N +++ +I++AHSRFPVI EDKD + GIL AKDLL +M +
Sbjct: 70 RDIMIPRSQMITLKRNLQTLDECLDVIESAHSRFPVISEDKDHIIEGILMAKDLLPFMRSD 129

35 Query: 120 PEQFHLKSILRPVAVFVPEGKSLTALLKEFRQNRHMAIVIDEYGGTSGLVTFEDIIEQIV 179
E F + +LR AV VPE K + +LKEFR QR HMAIVIDE+GG SGLVT EDI+E IV
Sbjct: 130 AEAFSMDKVLRLQAVVVPESKRVDRLMKEFRSQRYHMAIVIDEFGGVSGLVTTIEDILELIV 189

Query: 180 GDIEDEFDEDESADNIHAVSAERWRIHAATEIEDINAFFGTEYSSEEADT 229
G+IEDE+DE++ D +S W + A IED N FGT +S EE DT
Sbjct: 190 GEIEDEYDEEDDID-FRQLSRHTWTVRALASIEDFNEAFGTHFSDEEVD 238

40 Based on this analysis, including the amino acid homology to the TlyC hemolysin-homologue from
H. influenzae (hemolysins are secreted proteins), it was predicted that the proteins from
N.meningitidis and *N.gonorrhoeae* are secreted and could thus be useful antigens for vaccines or
diagnostics.

ORF5-1 (30.7kDa) was cloned in the pGex vector and expressed in *E.coli*, as described above. The
45 products of protein expression and purification were analyzed by SDS-PAGE. Figure 2A shows
the results of affinity purification of the GST-fusion protein. Purified GST-fusion protein was used
to immunise mice, whose sera were used for Western blot analysis (Figure 1B). These experiments
confirm that ORF5-1 is a surface-exposed protein, and that it is a useful immunogen.

Example 5

50 The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 29>:

1 ATGCGCGGCG GCAGGCCGGA TTCCGTTACC GTGCAGATTA TCGAAGGTTTC
51 GCGTTTTTCG CATATGAGGA AAGTCATCGA CGCAACGCCG GACATCGGAC

101 ACGACACCAA AGGCTGGAGC AATGAAAAAC TGATGGCGGA AGTTGCGCCC
 151 GATGCCTTCA GCGGCAATCC TGAAGGGCAG TTTTCCCCG ACAGCTACGA
 201 AATCGATGCG GCGGCGAGTG ATTTGCAGAT TTACCAAACC GCCTACAAGG
 251 GCATGCAAC GCCGCCTGAA TGAAGGCATG GGAAAGCAGG CAGGACGGGC
 5 301 TGCCTTATAA AAACCTTAT GAAATGCTGA TTATGGCGAr CCTGGTCGAA
 351 AAGGAAACAG GGCATGAAGC CGAsCsCGAC CATGTcGGCTT CCGTCTTCGT
 401 CAACCGCCTG AAAATCGGTA TCGCCTGCA AACCGAssCG TCCGTGATTT
 451 ACGGCATGGG TCGGCATAC AAGGGCAAAA TCCGTAAAGC CGACCTGCGC
 501 CGCGACACGC CGTACAACAC CTACACGCGC GCGGGTCTGC CGCAACCCC
 10 551 GATTGCGCTG CCC..

This corresponds to the amino acid sequence <SEQ ID 30; ORF7>:

1 MRGGRPDsvT VQIIeGSRfS HMRKVIDATP DIGHDTKGWS NEKLMAEVAP
 51 DAFSGNPEGQ FFPDSYEIDA GGSdlQIYQT AYKAMQRRLN EAWESRODGL
 101 PYKNPYEMLI MAXLVEKETG HEAXXDHVAS VFVNRlKIGM RLQTXSVIY
 15 151 GMAAYKGKI RKADLRRDTP YNTYTRGGLP PTPIALP..

Further sequence analysis revealed the complete DNA sequence <SEQ ID 31>:

1 ATGTTGAGAA AATGTTGAA ATGGTCTGCC GTTTTTTTGA CCGTGTCGGC
 51 AGCCGTTTTTC GCCGCGCTGC TTTTGTTC TAAGGATAAC GGCAGGGCAT
 101 ACCGAATCAA AATGCCAAA AACCAGGTA TTTCTCGGT CGGCAGGAAA
 151 CTTGCCGAAG ACCGCATCGT GTTCAGCAGG CATGTTTTGA CGGCGGCGGC
 20 201 CTACGTTTTG GGTGTGCACA ACAGGCTGCA TACGGGACG TACAGATTGC
 251 CTTCGGAAGT GTCTGCTTGG GATATCTTGC AGAAAATGCG CGGCGGACAG
 301 CCGGATTCCG TTACCGTGCA GATTATCGAA GGTTCGCGTT TTTGCGATAT
 351 GAGGAAAGTC ATCGACGCAA CGCCGACAT CGGACACGAC ACCAAAGGCT
 25 401 GGAGCAATGA AAAACTGATG GCGGAAGTTG CGCCGATGC CTTACGGCG
 451 AATCCTGAAG GGCAGTTTTT CCGGACAGC TACGAAATCG ATGCGGGCGG
 501 CAGTGATTTG CAGATTTACC AAACCGCCTA CAAGGCGATG CAACGCGGCC
 551 TGAATGAGGC ATGGGAAAGC AGGCAGGACG GGCTGCCTTA TAAAAACCT
 601 TATGAAATGC TGATTATGGC GAGCCTGGTC GAAAAGGAAA CAGGGCATGA
 30 651 AGCCGACCGC GACCATGTCG CTTCCGTCTT CGTCAACCGC CTGAAAATCG
 701 GTATGCGCCT GCAAACCGAC CCGTCCGTGA TTTACGGCAT GGTGCGGCA
 751 TACAAGGGCA AAATCCGTAA AGCCGACCTG CGCCGCGACA CGCCGTACAA
 801 CACCTACAG CGCGGCGGTC TGCCGCCAAC CCGATTGCG CTGCCCGGCA
 851 AGGCGGCACT CGATGCCGCC GCCCATCCGT CCGGCGAAAA ATACCTGTAT
 35 901 TTCGTGTCCA AAATGGACGG CACGGGCTTG AGCCAGTTCA GCCATGATTT
 951 GACCGAACAC AATGCCGCCG TCCGCAATA TATTTTGAAA AAATAA

This corresponds to the amino acid sequence <SEQ ID 32; ORF7-1>:

1 MLRKLLKWSA VFLTVSAAVF AALLFVPKDN GRAYRIKIAK NQGISSVGRK
 40 51 LAEDRIVFSR HVLTAAYVL GVHNRlHTGT YRLPSEVS AW DILQKMRGR
 101 PDSVTVQIIE GSrFSHMRKV IDATPDIGHD TKGWSNEKIM AEVAPDAFSG
 151 NPEGQFFPDS YEIDAGGSDl QIYQTAYKAM QRRLNEAWES RQDGLPYKNP
 201 YEMLIMASLV EKETGHEADR DHVASVFVNR LKIGMRLQTD PSVIYGMGAA
 251 YEKIRKADL RRDTPYNTYT RGLPPTPIA LPGKAALDAA AHPSGEKYLY
 301 FVSKMDGTGL SQFSHDlTEH NAAVRKYILK K*

45 Computer analysis of this amino acid sequence gave the following results:

Homology with hypothetical protein encoded by yceg gene (accession P44270) of *H. influenzae*

ORF7 and yceg proteins show 44% aa identity in 192 aa overlap:

ORF7 1 MRGGRPDsvTVQIIeGSRfSHMRKVIDATPDIGHDTKGWSNEKLMA-----EVAPDAFSG 55
 + G+ V+ IEG F RK ++ P + K SNE++ A ++ +
 50 yceg 102 LNSGKEVQFNWKWIEGKTFKDWKDLNAPHLVQTLKDKSNEEIFALLDLDPIDIGONLELK 161
 ORF7 56 NPEGQFFPDSYEIDAGGSDlQIYQTAYKAMQRRLNEAWESRQDGLPYKNPYEMLI MAXLV 115
 N EG +PD+Y +DL++ + + + M++ LN+AW R + LP NPYEMLI+A +V
 55 yceg 162 NVEGWLYPDYNTYPKSTDLLELLKRSaERMKKALNKAWNERDEDLPLANPYEMLILASIV 221
 ORF7 116 EKETGHEAXXDHVASVFVNRlKIGMRLQTXSVIYGMGAAYKGKIRKADLRRDTPYNTYT 175
 EKETG VASVF+NRlK M+LQT +VIYGMG Y G IRK DL TPNNTY
 yceg 222 EKETGIANERAKVASVFINRLKAKMKLQTDPTVIYGMGENYNGNIRKDLTKTPYNTYV 281

ORF7 176 RGGLPPTPIALP 187
GLPPTPIA+P
yceg 282 IDGLPPTPIAMP 293

The complete length YCEG protein has sequence:

```

5      1 MKKFLIAILL LILILAGVAS FSYYKMTEFV KTPVNVQADE LLTIERGTTS
      51 SKLATLFEQE KLIADGKLLP YLLKLKPELN KIKAGTYSLE NVKTVQDLLD
     101 LLNSGKEVQF NVKWIEGKTF KDWRKDLENA PHLVQTLKDK SNEEIFALLD
     151 LPDIGQNLEL KNVEGWLYPD TYNYPKSTD LELLKRSAER MKKALNKAWN
     201 ERDEDLPLAN PYEMLILASI VEKETGIANE RAKVASVFIN RLKAKMKLQT
     251 DPTVIYGMGE NYNGNIRKGD LETKTPYNTY VIDGLPPTPI AMPSESSLQA
     301 VANPEKTDYF YFVADGSGGH KFTRNLNEHN KAVQEYLRWY RSQKNAK

```

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF7 shows 95.2% identity over a 187aa overlap with an ORF (ORF7a) from strain A of *N.*

15 *meningitidis*:

```

                                     10      20      30
orf7.pep                               MRGGRPDSVTVQIIIEGSRFSHMRKVIDATP
                                     |||
orf7a      AAYVLGVHNRRLHTGYRLPSEVSAWDILQKMRGGRPDSVTVQIIIEGSRFSHMRKVIDATP
      70      80      90      100     110     120

                                     40      50      60      70      80      90
orf7.pep      DIGHDTKGWSNEKLMAEVAPDAFSGNPEGQFFPDSEIDAGGSDLQIYQTAYKAMQRRLN
      |||
orf7a      DIEHDTKGWSNEKLMAEVAPDAFSGNPEGQFFPDSEIDAGGSDLRIYQIAYKAMQRRLN
      130     140     150     160     170     180

                                     100     110     120     130     140     150
orf7.pep      EAWESRQDGLPYKNPYEMLIMAXLVEKETGHEAXDXHVASVFNRLKIGMRLQTXSVIY
      |||
orf7a      EAWESRQDGLPYKNPYEMLIMASLIEKETGHEADRDHVASVFNRLKIGMRLQTDPSVIY
      190     200     210     220     230     240

                                     160     170     180
orf7.pep      GMGAAYKGKIRKADLRRDTPYNTYTRGGLPPTPIALP
      |||
orf7a      GMGAAYKGKIRKADLRRDTPYNTYTRGGLPPTPIALPGKAALDAAHPSGEKYLYFVSKM
      250     260     270     280     290     300

orf7a      DGTGLSQFSHDLTEHNAAVRKYILKKX
      310     320     330

```

The complete length ORF7a nucleotide sequence <SEQ ID 33> is:

```

45      1 ATGTTGAGAA AATTGTTGAA ATGGTCTGCC GTTTTTTTGA CCGTATCGGC
      51 AGCCGTTTTC GCCGCGCTGC TTTTCGTCCC TAAAGACAAC GGCAGGGCAT
     101 ACAGGATTAA AATTGCCAAA AACCAGGGTA TTTTCGTCGGT CGGCAGGAAA
     151 CTTGCCGAAG ACCGCATCGT GTTCAGCAGG CATGTTTTGA CGGCGGCGGC
     201 CTACGTTTTG GGTGTGCACA ACAGGCTGCA TACGGGGACG TACAGACTGC
     251 CTTCGGAAGT GTCTGCTTGG GATATCTTGC AGAAAATGCG CGGCGGCAGG
     301 CCGGATTCCG TTACCGTGCA GATTATCGAA GGTTCCGCTT TTTTCGCATAT
     351 GAGGAAAGTC ATCGACGCAA CGCCCACAT CGAACACGAC ACCAAAGGCT
     401 GGAGCAATGA AAAACTGATG GCGGAAGTTG CCCCTGATGC CTTACAGCGC
     451 AATCCTGAAG GGCAGTTTTT CCCCACAGC TACGAAATCG ATGCGGCGCG
     501 CAGCGATTTA CGGATTTACC AAATCGCCTA CAAGGCGATG CAACGCCGAC
     551 TGAATGAGGC ATGGGAAAGC AGGCAGGACG GGCTGCCTTA TAAAAACCCT
     601 TATGAAATGC TGATTATGGC GAGCCTGATC GAAAAGGAAA CAGGGCATGA
     651 AGCCGACCGC GACCATGTCG CTTCCGTCTT CGTCAACCGC CTGAAAATCG
     701 GTATGCGCCT GCAAACCGAC CCGTCCGTGA TTTACGGCAT GGGTGC GGCA
     751 TACAAGGGCA AAATCCGTAA AGCCGACCTG CGCCGCGACA CGCCGTACAA
     801 CACCTACACG CGCGGCGGTC TGCCGCCAAC CCCGATCGCG CTGCCCGGCA
     851 AGGCGGCACT CGATGCGGCC GCCATCCGT CCGGTGAAAA ATACCTGTAT
     901 TTCGTGTCCA AAATGGACGG TACGGGCTTG AGCCAGTTCA GCCATGATTT
     951 GACCGAACAC AACCGCGCCG TTCGCAATA TATTTTGAAA AAATAA

```

This is predicted to encode a protein having amino acid sequence <SEQ ID 34>:

```

      1 MLRKLLKWSA VFLTVSAAVF AALLFVPKDN GRAYRIKIAK NQGISSVGRK
     51 LAEDRIVFSR HVLTAAYVL GVHNRLHTGT YRLPSEVSAW DILQKMRGGR
    101 PDSVTVQIIE GSRFSHMRKV IDATPDIEHD TKGWSNEKLM AEVAPDAFSG
5     151 NPEGQFFPDS YEIDAGGSDL RIYQIAYKAM QRRLEAWES RQDGLPYKNP
    201 YEMLIMASLI EKETGHEADR DHVASVFVNR LKIGMRLQTD PSVIYGMGAA
    251 YKGKIRKADL RRDTPYNTYT RGGLPPTPIA LPGKAALDAA AHPSGEKYLY
    301 FVSKMDGTGL SQFSDLTEH NAAVRKYILK K*

```

A leader peptide is underlined.

10 ORF7a and ORF7-1 show 98.8% identity in 331 aa overlap:

```

      10      20      30      40      50      60
orf7a.pep MLRKLLKWSAVFLTVSAAVFAALLFVPKDNGRAYRIKIAKNQGISSVGRKLAEDRIVFSR
      |||
orf7-1    MLRKLLKWSAVFLTVSAAVFAALLFVPKDNGRAYRIKIAKNQGISSVGRKLAEDRIVFSR
      |||
15      10      20      30      40      50      60

      70      80      90     100     110     120
orf7a.pep HVLTAAYVLGVHNRLHTGT YRLPSEVSAWDILQKMRGGRPDSVTVQIIEGSRFSHMRKV
      |||
20      70      80      90     100     110     120
orf7-1    HVLTAAYVLGVHNRLHTGT YRLPSEVSAWDILQKMRGGRPDSVTVQIIEGSRFSHMRKV
      |||

      130     140     150     160     170     180
orf7a.pep IDATPDIEHDTKGWSNEKLM AEVAPDAFSGNPEGQFFPDSYEIDAGGSDLRIYQIAYKAM
      |||
25      130     140     150     160     170     180
orf7-1    IDATPDIGHDTKGWSNEKLM AEVAPDAFSGNPEGQFFPDSYEIDAGGSDLQIYQTAYKAM
      |||

      190     200     210     220     230     240
orf7a.pep QRRLEAWESRQDGLPYKNPYEMLIMASLIEKETGHEADRDHVASVFVNR LKIGMRLQTD
      |||
30      190     200     210     220     230     240
orf7-1    QRRLEAWESRQDGLPYKNPYEMLIMASLVEKETGHEADRDHVASVFVNR LKIGMRLQTD
      |||

      250     260     270     280     290     300
orf7a.pep PSVIYGMGAAYKGKIRKADLRRDTPYNTYTRGGLPPTPIALPGKAALDAAAHPSGEKYLY
      |||
35      250     260     270     280     290     300
orf7-1    PSVIYGMGAAYKGKIRKADLRRDTPYNTYTRGGLPPTPIALPGKAALDAAAHPSGEKYLY
      |||

      310     320     330
orf7a.pep FVSKMDGTGLSQFSDLTEHNAAVRKYILKKX
      |||
40      310     320     330
orf7-1    FVSKMDGTGLSQFSDLTEHNAAVRKYILKKX
      |||
45      310     320     330

```

Homology with a predicted ORF from *N.gonorrhoeae*

ORF7 shows 94.7% identity over a 187aa overlap with a predicted ORF (ORF7.ng) from *N. gonorrhoeae*:

```

50      orf7      MRGGRPDSVTVQIIEGSRFSHMRKV IDATPDIGHDTKGWSNEKLM AEVAPDAFSGNPEGQ      60
      orf7ng     MRGGRPDSVTVQIIEGSRFSHMRKV IDATPDIGHDTKGWSNEKLM AEVAPDAFSGNPEGQ      60

      orf7      FFPDSYEIDAGGSDLQIYQTAYKAMQRRLEAWESRQDGLPYKNPYEMLIMAXLVEKETG      120
55      orf7ng     FFPDSYEIDAGGSDLQIYQTAYKAMQRRLEAWAGRQDGLPYKNPYEMLIMASLIEKETG      120

      orf7      HEAXXDHVASVFVNR LKIGMRLQTXSVIYGMGAAYKGKIRKADLRRDTPYNTYTRGGLP      180
60      orf7ng     HEADRDHVASVFVNR LKIGMRLQDPSVIYGMGAAYKGKIRKADLRRDTPYNTYTGGLP      180

      orf7      PTPIALP

```

|| ||||
orf7ng PTRIALPGKAAMDAAAHPSGEKYLYFVSKMDGTGLSQFSHDLTEHNAAVRKYILKK 236

An ORF7ng nucleotide sequence <SEQ ID 35> is predicted to encode a protein having amino acid sequence <SEQ ID 36>:

5 1 MRGGRPDSVT VQIIEGSRFS HMRKVIDATP DIGHDTKGWS NEKLMAEVAP
51 DAFSGNPEGQ FFPDSYEIDA GGSDLQIYQT AYKAMQRRNL EAWAGRDGL
101 PYKNPYEMLI MASLIEKETG HEADRDHVAS VFNRLKIGM RLQTDPSVIY
151 GMGAAYKGI RKADLRDTP YNTYTGGGLP PTRIALPGKA AMDAAHPSG
201 EKLYFVSKM DGTGLSQFSH DLTEHNAAVR KYILKK*

10 Further sequence analysis revealed a partial DNA sequence of ORF7ng <SEQ ID 37>:

1 ..taccgaatca AGATTGCCAA AAATCAGGGT ATTTCGTCGG TCGGCAGGAA
51 ACTTGCCgaA GACCGCATCG TGTCAGCAG GCATGTTTTC ACAGCGGCGG
101 CCTACGTTT GGGTGTGCAC AACAGGCTGC ATACGGGGAC gTACAGATTG
151 CCTTCGGAAG TGTCTGCTTG GGATATCTTG CAGAAAATGC GCGGCGGCAG
201 GCCGGATTCC GTTACCGTGC AGATTATCGA AGTTTCGGCT TTTCGCATA
251 TGAGGAAAGT CATCGACGCA ACGCCCGACA TCGGACACGA CACCAAAGGC
301 TGGAGCAATG AAAAAGTATG GCGGGAAGTT GCGCCCGATG CCTTCAGCGG
351 CAATCCTGAA GGGCAGTTTT TTCCCGACAG CTACGAAATC GATGCGGGCG
401 GCAGCGATTG GCAGATTAC CAAACCGCT ACAAGGCGAT GCAACGCCGC
20 451 CTGAACGAGG CATGGGCAGG CAGGCAGGAC GGGCTGCCTT ATAAAAACCC
501 TTATGAAATG CTGATTATGG CGAGCCTGAT CGAAAAGGAA ACGGGGCATG
551 AGGCCGACCG CGACCATGTC GCTCCGTCT TCGTCAACCG CCTGAAATC
601 GGTATGCGCC TGCAAACCGA CCGTCCGTG ATTTACGCA TGGGTGCGGC
651 ATACAAGGC AAAATCCGTA AAGCCGACCT GCGCCGCGAC ACGCCGTACA
25 701 aCAccTatac gggcgggggc ttgccgcaa cccgattgc gctgccggc
751 Aaggcgcaa tggatgccgc cggccaccgc tccggcgaa aatacctgTa
801 ttctgtgtc AAAATGGACG GCACGGGCTT GAGCCAGTTC AGCCATGATT
851 TGACCGAACA CAACGCCGc gTcCGCAAT ATATTTTGAA AAAATAA

This corresponds to the amino acid sequence <SEQ ID 38; ORF7ng-1>:

30 1 ..YRIKIAKNQG ISSVGRKLAE DRIVFSRHVL TAAAYVLGVH NRLHTGTYRL
51 PSEVSAWDIL QKMRGGRPDS VTVQIIEGSR FSHMRKVIDA TPDIGHDTKG
101 WSNEKLMAEV APDAFSGNPE GQFFPDSYEI DAGGSDLQIY QTAYKAMQRR
151 LNEAWAGRD GLPYKNPYEM LIMASLIEKE TGHEADRDHV ASVFNRLKI
201 GMRLQTDPSV IYGMGAAYKG KIRKADLRD TPNTYTGGG LPPTRIALPG
35 251 KAAMDAAHP SGEKYLYFVS KMDGTGLSQF SHDLTEHNAA VRKYILKK*

ORF7ng-1 and ORF7-1 show 98.0% identity in 298 aa overlap:

	10	20	30	40	50	60
orf7-1.pep	KLLKWSAVFLTVSAAVFAALLFVPKDNGRAYRIKIAKNQGISSVGRKLAEDRIVFSRHVL					
40 orf7ng-1	YRIKIAKNQGISSVGRKLAEDRIVFSRHVL					
	70	80	90	100	110	120
orf7-1.pep	TAAAYVLGVHNRHLHTGTYRLPSEVSAWDILQKMRGGRPDSVTVQIIEGSRFSHMRKVIDA					
45 orf7ng-1	TAAAYVLGVHNRHLHTGTYRLPSEVSAWDILQKMRGGRPDSVTVQIIEGSRFSHMRKVIDA					
	130	140	150	160	170	180
orf7-1.pep	TPDIGHDTKGWSNEKLMAEVAPDAFSGNPEGQFFPDSYEIDAGGSDLQIYQTAYKAMQRR					
50 orf7ng-1	TPDIGHDTKGWSNEKLMAEVAPDAFSGNPEGQFFPDSYEIDAGGSDLQIYQTAYKAMQRR					
	100	110	120	130	140	150
55 orf7-1.pep	LNEAWESRQDGLPYKNPYEMLIMASLVEKETGHEADRDHVASVFNRLKIGMRLQTDPSV					
orf7ng-1	LNEAWAGRDGLPYKNPYEMLIMASLIEKETGHEADRDHVASVFNRLKIGMRLQTDPSV					
	160	170	180	190	200	210
60 orf7-1.pep	IYGMGAAYKGKIRKADLRDTPYNTYTGGGLPPTPIALPGKAALDAAHPSGEKYLYFVS					

-78-

```

      |||
orf7ng-1  IYGMGAAYKGKIRKADLRDTPYNTYTGGLFPTRIALPGKAAMDAAAHPSGEKYLIFVS
              220      230      240      250      260      270

5      310      320      330
orf7-1.pep  KMDGTGLSQFSHDLTEHNAAVRKYLKXX
              |||
orf7ng-1  KMDGTGLSQFSHDLTEHNAAVRKYLKXX
              280      290
10

```

In addition, ORF7ng-1 shows significant homology with a hypothetical *E. coli* protein:

```

15  sp|P28306|YCEG_ECOLI HYPOTHETICAL 38.2 KD PROTEIN IN PABC-HOLB INTERGENIC REGION
    gi|1787339 (AE000210) o340; 100% identical to fragment YCEG_ECOLI SW: P28306 but
    has 97 additional C-terminal residues [Escherichia coli] Length = 340
    Score = 79 (36.2 bits), Expect = 5.0e-57, Sum P(2) = 5.0e-57
    Identities = 20/87 (22%), Positives = 40/87 (45%)

20  Query:   10 GISSVGRKLAEDRIVFSRHLVTAAYVLGVHNRLLHTGTYRLPSEVSAWDILQKMRGGRPD 69
    G ++G +L D+I+ V + + GTYR +++ ++L+ + G+
    Sbjct:   49 GRLALGEQLYADKIINRPVRFQWLLRIEPLDSHFAGTYRFTPMQTVREMLKLLESGKEA 108

    Query:   70 SVTVQIIIEGSRFSHMRKVIDATPDIGH 96
    ++++EG R S K + P I H
    Sbjct:   109 QFPLRLVEGMRLSDYKQLREAPYIKH 135

25  Score = 438 (200.7 bits), Expect = 5.0e-57, Sum P(2) = 5.0e-57
    Identities = 84/155 (54%), Positives = 111/155 (71%)

30  Query:   120 EGQFFPDSYEIDAGGSDLQIYQTAYKAMQRRLEAWAGRQDGLPYKNPYEMLIMASLIEK 179
    EG F+PD++ A +D+ + + A+K M + ++ AW GR DGLPYK+ +++ MAS+IEK
    Sbjct:   158 EGFWFDPDTWMTANTTDVALLKRAHKMKVKAVDSAWEGRADGLPYKDKNLVTMASIIEK 217

    Query:   180 ETGHEADRDHVASVFVNRLKIGMRLQTDPSVIYGMGAAYKGKIRKADLRDTPYNTYTG 239
    ET ++RD VASVF+NRL+IGMRLQTD+VIYGMG Y GK+ +ADL T YNTYT
    Sbjct:   218 ETAVASERDKVASVFVNRLRIGMRLQTDPTVIYGMGERYNGKLSRADLETPTAYNTYTIT 277

35  Query:   240 GLPPTRIALPGKAAMDAAAHPSGEKYLIFVSKMDG 274
    GLPP IA PG ++ AAAHP+ YLYFV+ G
    Sbjct:   278 GLPPGAIATPGADSLKAAAHPAKTPYLYFVADGKG 312

40

```

Based on this analysis, including the fact that the *H. influenzae* YCEG protein possesses a possible leader sequence, it is predicted that the proteins from *N. meningitidis* and *N. gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 6

45 The following partial DNA sequence was identified in *N. meningitidis* <SEQ ID 39>:

```

50  1  CGTTTCAAAA TGTAACTGT GTTGACGGCA ACCTTGATTG CCGGACAGGT
    51  ATCTGCCGCC GGAGCGGTG CGGGGATAT GAAACAGCCG AAGCAAGTCG
    101 GAAAGGTTT CAGAAAGCAG CAGCGTTACA GCGAGGAAGA AATCAAAAAC
    151 GAACGCGCAC GGCTTGCGGC AGTGGCGGAG CCGGTTAATC AGATATTTAC
    201 GTTGCTGGGA GGGGAAACCG CCTTGCAAAA GGGGCAGGCG GGAACGGCTC
    251 TGGCAACCTA TATGCTGATG TTGGAACGCA CAAATCCCC CGAAGTCGCG
    301 GAACGCGCCT TGGAAATGGC CGTGTGCTG AACGCGTTG AACAGGCGGA
    351 AATGATTTAT CAGAAATGGC GGCAGATTGA GCCTATACCG GGTAAAGGCGC
    401 AAAAACGGGC GGGGTGGCTG CGGAACGTGC TGAGGGAAAG AGGAAATCAG
55  451 CATCTGGACG GACGGGAAGA AGTGCTGGCT CAGGCGGACG AAGGACAG

```

This corresponds to the amino acid sequence <SEQ ID 40; ORF9>:

```

    1  .RFKMLTVLTA TLIAGQVSAA GGGAGDMKQP KEVGKVERKQ QRYSEEEIKN
    51  ERARLAAVGE RVNQIFTLG GETALQKGQA GTALATYMLM LERTKSPEVA
    101  ERALEMAVSL NAFEQAEMIV QKWRQIEPIP GKAQKRAGWL RNVLRERGNQ

```

	1	ATGTTACCTA	ACCGTTTCAA	AATGTTAACT	GTGTTGACGG	CAACCTTGAT
5	51	TGCCGGACAG	GTATCTGCCG	CCGGAGGCGG	TGCCGGGGAT	ATGAAACAGC
	101	CGAAGGAAGT	CGGAAAGGTT	TTCAGAAAGC	AGCAGCGTTA	CAGCGAGGAA
	151	GAATCAAAA	ACGAAACGCG	ACGGCTTGCG	GCAGTGGCGG	AGCGGGTTAA
	201	TCAGATATTT	ACGTTGCTGG	GAGGGGAAAC	CGCCTTGCAA	AAGGGGCAGG
10	251	CGGGAACGCG	TCTGGCAACC	TATATGCTGA	TGTTGGAACG	CACAAAATCC
	301	CCCGAAGTCG	CGGAACGCGC	CTTGAAATGC	CGCGTGTCCG	TGAACCGGTT
	351	TGAACAGCGG	GAAATGATTT	ATCAGAAATG	GCGGCAGATT	GAGCCTATAC
	401	CGGTAAGGC	GCAAAAACGG	GCGGGGTGGC	TGCGGAAACG	GCTGAGGGAA
15	451	AGAGGAAATC	AGCATCTGGA	CGGACTGGAA	GAAGTGCTGG	CCTCAGCGGA
	501	CGAAGGACAG	AACCGCAGGG	TGTTTTTATT	TTGGGCACAA	CGCGCGGTGC
	551	AACAGGACGG	GTTGGCGCAA	AAAGCATCGA	AAGCGGTTCC	CCGCGCGGCG
	601	TTGAAATATG	AACATCTGCC	CGAAGCGGCG	GTTGCCGATG	TGGTGTTTCA
20	651	CGTACAGGGA	CGCGAAAAGG	AAAAGGCAAT	CGGAGCTTTG	CAGCGCTTGG
	701	CGAAGCTCGA	TACGAAATA	TTGCCCCCCA	CTTTAATGAC	TGTGCGTCTG
	751	ACTGCACGCA	AATATCCCGA	AATACTCGAC	GGCTTTTTTC	AGCAGACAGA
	801	CACCCAAAAC	CTTTTCGGCG	TCTGGCAGGA	AATGGAATTT	ATGAATCTGG
25	851	TTTCCCTGCA	CAGGCTGGAT	GATGCCTATG	CGCGTTTGAA	CGTGCCTGTT
	901	GAACGCAATC	CGAATCGAGA	CCTGTATATT	CAGGCAGGCA	TATTGGCGGC
	951	AAACCGAAAA	GAAGGTGCTT	CCGTATTCGA	CGGCTACGCC	GAAAAGGCGA
	1001	ACCGCAGGGG	CAGCGAGGAA	CAGCGGAGCA	GGCGCGCGCT	AACGGCGGCG
30	1051	ATGATGTATG	CCGACCGCAG	GGATTACGCC	AAAGTCAGGC	AGTGGCTGAA
	1101	AAAAGTATCC	GCGCCGGAAT	ACCTGTTTCA	CAAAGGTGTG	CTGGCGGCTG
	1151	CGGCGGCTGT	CGAGTTGGAC	GGCGGCAGGG	CGGCTTTGCG	GCAGATCGGC
	1201	AGGGTGCGGA	AATTTCCCGA	ACAGCAGGGG	CGGTATTTTA	CGGCAGACAA
35	1251	TTTGTCCAAA	ATACAGATGC	TCGCCCTGTC	GAAAGTCGCC	GATAAACGGG
	1301	AGGCTTTGAG	GGGGTTGGAC	AAGATTATCG	AAAAACCGCC	TGCCGGCAGT
	1351	AATACAGAGT	TACAGGCAGA	GGCATTTGTA	CAGCGGTTCAG	TTGTTTTACGA
	1401	TCGGCTTGCG	AAGCGGAAAA	AAATGATTTT	AGATCTTGAA	AGGGCGCTTC
40	1451	GGCTTGACCC	CAGTAACGCT	CAGATTATGA	ATAATCTGGG	CTACAGGCTG
	1501	CTGACCGATT	CCAAACGTTT	GGACGAAGGT	TTCGCCCTGC	TTTACAGCGGC
	1551	ATACCAAAATC	AAACCGGACG	ATACCGCTGT	CACACGACAG	ATAGGCTGGT
	1601	CGTATTACCT	GAAAGGCGAC	GCGGAAAGCG	CGCTGCGGTA	TGTGCGGTAT
45	1651	TCGTTTGAAA	ACGACCCCGA	GCCCCAAGTT	GCCGCCCATT	TGGGCGAAGT
	1701	GTTGTGGGCA	TTGGGCGAAC	GCGATCAGCG	GGTTGACGTA	TGGACGCGAGG
	1751	CGGCACACCT	TACGGGAGAC	AAGAAAAATAT	GGCGGGAAAC	GCTCAACCTT
	1801	CACGGCATTC	CATTGCCCCA	ACCTTCCCAG	AAACCTCGGA	AATAA

	1	MLPNRFKMLT	VLATATLIAGQ	VSAAGGGAGD	MKQPKEVGKV	FRKQQRYSSE
	51	EIKNERARLA	AVGERVNIQIF	TLLGGETALQ	KGOAGTALAT	YMLMLERTKS
	101	PEVAERALEM	AVSLNAFEQA	EMIQKWRQI	EPIPGKAQKR	AGWLRNVLRE
45	151	RGNQHLDDLE	EVLAQADEGQ	NRRVFLLLAQ	AAVQDGLAQ	KASKAVRRAA
	201	LKYEHLFEAA	VDVVFSVQG	REKEKAIGAL	QRLAKLDEI	LPPPTLMTLRL
	251	TARKYPEILD	GFFEQTDTQN	LSAVWQEME	MNLVSLHRLD	DAYARLNVLL
	301	ERNPNADLYI	QAAILAANRK	EGASVIDGYA	EKAYGRGTEE	QRSRAALTAQ
	351	MMYADRDRYA	KVRQWLKKVS	APEYLFDKGV	LAAAAAVELD	GGRAALRIGI
	401	RVRKLPEQQG	RYFTADNLKS	IQLMALSKLP	DKREALRGD	KIIEKPPAGS
50	451	NTELQAEALV	QRSVVYDRIG	KRKKMISDLE	RAFRLAPDNA	QIMNNLGYSL
	501	LTDSCRLEDE	FALLQTAYQI	NPDDTAVNDS	TGWAYYLKGD	AESALPYLRY
	551	SFENDPEPEV	AAHLGEVLWA	LGERDQAVDV	WTQAAHLTGD	KKIWRETLKR
	601	HGIALPQPSR	KPRK*			

55 Homology with a predicted ORF from *N.meningitidis* (strain A)

```

                                10      20      30      40      50
60 orf9.pep    RFKMLT1TVLTATLIAGQVSAAGGGGADMKQPKEVGKVFRKQORYSEEEIKNERARLA
              ||::||:|:|:||||: | | |:| | | | | | | | | | | | | | | | | | | |
orf9a        MLPARETILSVLAAALLAGOAYAA--GAADAKPPKEVGKVFRKQORYSEEEIKNERARLA

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-80-

		10	20	30	40	50	
		60	70	80	90	100	110
5	orf9.pep	AVGERVNQIFTLGGETALQKGQAGTALATYMLMLERTKSPEVAERALEMAVSLNAFEQA					
	orf9a	AVGERVNQIFTLGXETALQKGQAGTALATYMLMLERTKSPEVAERALEMAVSLNAFEQA					
		60	70	80	90	100	110
10	orf9.pep	EMIQKWRQIEPIPGKAQKRAGWLRNVLRERGNQHLDGREEVLAQADEGQ					
	orf9a	EMIQKWRQIEPIPGKAQKRAGWLRNVLRERGNQHLDGLEEXLAQADEXQNRVFLLLAQ					
		120	130	140	150	160	170
15	orf9a	AAVQQDGLAQKASKAVRRAALRYEHLPEAAVADVFSVOXREKEKAIGALQRLAKLDTEI					
		180	190	200	210	220	230

The complete length ORF9a nucleotide sequence <SEQ ID 43> is:

1	ATGTTACCCG	CCCGTTTCAC	CATTTTATCT	GTGCTCGCGG	CAGCCCTGCT
51	TGCCGGGCAG	GCGTATGCCG	CCGGCGCGCG	GGATGCGAAG	CCGCCGAAGG
101	AAGTCGGAAG	GGTTTTCAGA	AAGCAGCAGC	GTTACAGCGA	GGAAGAAATC
151	AAAAACGAAC	GCGCACGGCT	TGCGGCAGTG	GGCGAGCGGG	TTAATCAGAT
201	ATTTACGTTG	CTGGGANGGG	AAACCGCCTT	GCAAAAGGGG	CAGGCGGGAA
251	CGGCTCTGGC	AACCTATATG	CTGATGTTGG	AACGCACAAA	ATCCCCGAA
301	GTCGCCGAAC	GCGCCTTGGA	AATGGCCGTG	TCNCTGAACG	CGTTTGAACA
351	GGCGGAAATG	ATTTATCAGA	AATGGCGGCA	GATTGAGCCT	ATACCGGGTA
401	AGGCGCAAAA	ACGGGCGGGG	TGGCTGCGGA	ACGTGCTGAG	GGAAGAGGA
451	AATCAGCATC	TAGACGGACT	GGAAGAANTG	CTGGCTCAGG	CGGACGAANG
501	ACAGAACCGC	AGGGTGTGTT	TATTGTTGGC	ACAAGCCGCC	GTGCAACAGG
551	ACGGGTTGGC	GCAAAAAGCA	TCGAAAAGCG	TTCGCGCGCG	GGCGTTGAGA
601	TATGAACATC	TGCCCCAAGC	GGCGTTGCC	GATGTGGTGT	TCAGCGTACA
651	GGNACGCGAA	AAGGAAAAGG	CAATCGGAGC	TTTGACGCGT	TTGGCGAAGC
701	TCGATACGGA	AATATGCCCC	CCCACCTTAA	TGACGTTGCG	TCTGACTGCA
751	CGCAAAATATC	CCGAAATACT	CGACGGCTTT	TTCGAGCAGA	CAGACACCCA
801	AAACCTTTTC	CCCGTCTGGC	AGGAAATGGA	AATTATGAAT	CTGGTTTCCC
851	TGCACAGGCT	GGATGATGCC	TATGCGCGTT	TGAACGTGCT	GTTGGAACGC
901	AATCCGAATG	CAGACCTGTA	TATTGAGGCA	GCGATATTGG	CGGCAAAACCG
951	AAAAGAANGT	GCTTCCGTGA	TCGACGGCTA	CGCCGAAAAG	GCATACGGCA
1001	GGGGGACGGG	GGAACAGCGG	GGCAGGGCGG	CAATGACGGC	GGCGATGATA
1051	TATGCCGACC	GAAGGGATTA	CACCAAAGTC	AGGCAGTGGT	TGAAAAAAGT
1101	GTCCGCGCCG	GAATACCTGT	TCGACAAAAG	TGTGCTGGCG	GCTGCGGCGG
1151	CTGTCGAGTT	GGACNGCGGC	AGGGCGGCTT	TGCGGCAGAT	CGGCAGGGTG
1201	CGGAAACTTC	CCGAACAGCA	GGGGCGGTAT	TTTACGGCAG	ACAATTTGTC
1251	CAAAATACAG	ATGTTGCGCC	TGTCGAAGCT	GCCCGACAAA	CGGGAGGCTT
1301	TGAGGGGGTT	GGACAAGATT	ATCGAAAAAC	CGCCTGCCGG	CAGTAATACA
1351	GAGTTACAGG	CAGAGGCATT	GGTACAGCGG	TCAGTTGTTT	ACGATCGGCT
1401	TGGCAAGCGG	AAAAAATGA	TTTCAGATCT	TGAAAGGGCG	TTGAGGCTTG
1451	CACCCGATAA	CGCTCAGATT	ATGAATAATC	TGGGCTACAG	CCTGCTTTCC
1501	GATTCCAAAC	GTTTGACAGA	AGGCTTCGCC	CTGCTTCAGA	CGGCATACCA
1551	AATCAACCCG	GACGATACCG	CTGTCAACGA	CAGCATAGGC	TGGGCGTATT
1601	ACCTGAAANG	CGACGCGGAA	AGCGCGCTGC	CGTATCTGCG	GTATTGTTTT
1651	GAAAACGACC	CCGAGCCCGA	AGTTGCCGCC	CATTGCGGCG	AAGTGTGTGT
1701	GGCATTGGGC	GAACGCGATC	AGGCGGTTGA	CGTATGGACG	CAGGCGGCAC
1751	ACCTTACGGG	AGACAAGAAA	ATATGGCGGG	AAACGCTCAA	ACGTACCGGC
1801	ATCGCATTGC	CCCAACCTTC	CCGAAAACCT	CGGAAATAA	

55 This encodes a protein having amino acid sequence <SEQ ID 44>:

1	MLPARFTILS	VLAAALLAQ	AYAAGAADAK	PPKEVGKVFR	KQORYSEEEI
51	KNERARLAAV	GERVNQIFTL	LGXETALQKG	QAGTALATYM	LMMLERTKSPE
101	VAERALEMAV	SLNAFEQAEM	IYQKWRQIEP	IPGKAQKRAG	WLRNVLRERG
151	NQHLDGLEEX	LAQADEXQNR	RVFLLLAQAA	VQQDGLAQKA	SKAVRRAALR
201	YEHLPAAAVA	DVVFVSQXRE	KEKAIGALQR	LAKLDTEILP	PTLMTLRLTA
251	RKYPEILDGF	FEQTDTONLS	AVWQEMEIMN	LVSLHRLDDA	YARLNVLLER
301	NPNADLYIQA	AILAANRKEX	ASVIDGYAEK	AYGRGTGEQR	GRAAMTAAMI
351	YADRRDYTKV	RQWLKKVSAP	EYLFDKGVLA	AAAVELDXG	RAALRQIGRV
401	RKLPEQGRY	FTADNLSIQ	MFALSKLPDK	REALRGLDKI	IEKPPAGSNT
451	ELQAEALVQR	SVVYDRLGKR	KKMISDLERA	FRLAPDNAQI	MNNLGYSLLS
501	DSKRLDEGFA	LLQTAYQINP	DDTAVNDSIG	WAYYLLKXDAE	SALPYLRYSF
551	ENDPEPEVAA	HLGEVLWALG	ERDQAVDVWT	QAAHLTGDKK	IWRETLKRHG

601 IALPQPSRKPK RK*

ORF9a and ORF9-1 show 95.3% identity in 614 aa overlap:

5	orf9a.pep	10 20 30 40 50	MLPARFTILSVLAAALLAGQAYAAG--AADAKPPKEVGKVFRKQQRYSSEEEKNERARLA
	orf9-1	10 20 30 40 50 60	MLPNRFKMLTVLTATLIAGQVSAAGGGAGDMKQPKEVGKVFRKQQRYSSEEEKNERARLA
10	orf9a.pep	60 70 80 90 100 110	AVGERVNQIFITLLGXETALQKGQAGTALATYMLMLERTKSPEVAERALEMAVSLNAFEQA
	orf9-1	70 80 90 100 110 120	AVGERVNQIFITLLGGETALQKGQAGTALATYMLMLERTKSPEVAERALEMAVSLNAFEQA
15	orf9a.pep	120 130 140 150 160 170	EMIQKWRQIEPIPGKAQKRAGWLRNVLRERGNQHLDGLEEXLAQADEQNRRVFLLLAQ
	orf9-1	130 140 150 160 170 180	EMIQKWRQIEPIPGKAQKRAGWLRNVLRERGNQHLDGLEEVLAQADEQNRRVFLLLAQ
20	orf9a.pep	180 190 200 210 220 230	AAVQQDGLAQKASKAVRRAALRYEHLPEAAVADVVSQXREKEKAIGALQRLAKLDTEI
	orf9-1	190 200 210 220 230 240	AAVQQDGLAQKASKAVRRAALKYEHLPEAAVADVVSQGREKEKAIGALQRLAKLDTEI
25	orf9a.pep	240 250 260 270 280 290	LPPTLMTLRLTARKYPEILDGFFEQTDTQNLSAVWQEMEIMNLVSLHRLDDAYARLNVLL
	orf9-1	250 260 270 280 290 300	LPPTLMTLRLTARKYPEILDGFFEQTDTQNLSAVWQEMEIMNLVSLHRLDDAYARLNVLL
30	orf9a.pep	300 310 320 330 340 350	ERNPNADLYIQAAILAANRKEASVIDGYAEKAYGRGTGEQGRGAAMTAAMIYADRRDYT
	orf9-1	310 320 330 340 350 360	ERNPNADLYIQAAILAANRKEGASVIDGYAEKAYGRGTEEQRSRAALTAAMMYADRRDYA
35	orf9a.pep	360 370 380 390 400 410	KVRQWLKKVSAPEYLFDKGVLA AAAA VELDXGRAALRQIGRVKRLPEQQGRYFTADNLSK
	orf9-1	370 380 390 400 410 420	KVRQWLKKVSAPEYLFDKGVLA AAAA VELDGGRAALRQIGRVKRLPEQQGRYFTADNLSK
40	orf9a.pep	420 430 440 450 460 470	IQMFALSKLPDKREALRGLDKIIEKPPAGSNTTELQAEALVQRSVVYDRLGKRKKMISDLE
	orf9-1	430 440 450 460 470 480	IQMLALSKLPDKREALRGLDKIIEKPPAGSNTTELQAEALVQRSVVYDRLGKRKKMISDLE
45	orf9a.pep	480 490 500 510 520 530	RAFRLAPDNAQIMNNLGYSLLSDSKRLDEGFALLQTAYQINPDDTAVNDSIGWAYYLKXD
	orf9-1	490 500 510 520 530 540	RAFRLAPDNAQIMNNLGYSLLTDSKRLDEGFALLQTAYQINPDDTAVNDSIGWAYYLKGD
50	orf9a.pep	540 550 560 570 580 590	AESALPYLRYSFENDPEPEVA AHLGEVLWALGERDQAVDVWTQAAHLTGDKKIWRETLKR
	orf9-1	550 560 570 580 590 600	AESALPYLRYSFENDPEPEVA AHLGEVLWALGERDQAVDVWTQAAHLTGDKKIWRETLKR
55	orf9a.pep	600 610	HGIALPQPSRKPKRX
	orf9-1	610	HGIALPQPSRKPKRX

5	orf9	RFKMLTVLTATLIAGQVSAAGGGAGDMKQPKVEGVKVRFKQQRYSEEEIKNERAR	54
	orf9ng	MIMLPARFTILSVLAAALLAGQAYAA--GAADVLPKEGVKVLKRHHRYSEEEIKNERAR	58
10	orf9	LAAVGERVNQIFTLGGETALQKGQAGTALATYMLMLERTKSPEVAERALEMAVSLNAFE	114
	orf9ng	LAAVGERVNRVFTLLGGETALQKGQAGTALATYMLMLERTKSPEVAERALEMAVSLNAFE	118
	orf9	QAEMIYQKWRQIEPIPGKAQKRAGWLRNVLRRGNQHLDGREEVLQAADEGQ	166
	orf9ng	QAEMIYQKWRQIEPIGGEAQKPAGWLRNVLKEGGNPHLDRLLEEVAQSDYVHOPMIFLLL	178

20

1	MIMLPARFTI	LSVLAAALLA	GQAYAAGAAD	VELPKEVGKV	LRKHRRYSEE
51	EIKNERARLA	AVGERVNRVF	TLLGGETALQ	KKQAGATALT	YMLMLERTKS
101	PEVAERALEM	AVSLNAFEQA	EMIYQKWRQI	EPIPGEAQKP	AGWLNRVLKE
151	GGNPHLDRLE	EVPAQSDYVH	QPMIFLLLVQ	AAVQHGGVAA	KPSKAVRPA
201	YNYEVLPETA	GADAVFCVQG	PQYEKAIQSF	PPCGRNQTE	NIAPPFNELF
251	RPTARLSPK	LLORFFTEP	NLAKPFPPPG	PEMETYOTGF	PEPLTRNNPT

25 Further sequence analysis revealed the complete length ORF9ng DNA sequence <SEQ ID 47>:

	1	ATGTTACCCG	CCCGTTTCAC	TATTTTATCT	GTCCTCGCAG	CAGCCCTGCT
	51	TGCCCGACAG	GCGTATGCTG	CCGGCGCGGC	GGATGTGGAG	CTGCCGAAGG
	101	AAGTCGGAAA	GGTTTAAAG	AAACATCGGC	GTTACAGCGA	GGAAGAAATC
30	151	AAAAACGAAC	GCGCACGGCT	TGCGGCAGTG	GTCACACGGG	TCAACAGGGT
	201	GTTTACGCTG	TTGGGCGGTG	AAACGGCTTT	GCAGAAAGGG	CAGGCGGGAA
	251	CGGCTCTGGC	AACCTATATG	CTGATGTTGG	AACGCACAAA	ATCCCCCGAA
	301	GTCGCCGAAC	GCGCCTTGGA	AATGGCCGTG	TCGCTGAACG	CGTTTGAACA
	351	GGCGGAAATG	ATTTATCAGA	AATGgcggca	gatcgagcct	ataCcggggtg
35	401	aggcgcaaaa	accgGcgggG	tggctgcgga	acgtattgaa	ggaagggGga
	451	aaTCAGCATC	TGGAcggggt	gaaagaggTG	CtgggcgaAT	cggacgatGT
	501	GCAAAAacgc	aggaTATTTT	TGCTGTGGT	GCAAGCCGCC	GTGCagcagg
	551	gTGGGGTGGC	TCAAAAAGCA	TCGAAAGCGG	TTCGCcgtgc	GGcgttgaAG
	601	TATGAACATC	TGCCcgaagc	ggcggTGGCC	GATGcggTGT	TCGGCGTACA
40	651	GGGACGCGAA	AAGGAAAagg	caaTCGAAGC	TTTGACAGCT	TTGGCGAAGC
	701	TCGATACGGA	AATATTGCC	CCCACTTTAA	TGACGTTGCG	TCTGACTGCA
	751	CGCAAATATC	CCGAAATACT	CGACGGCTTT	TTCGAGCAGA	CAGACACCCA
	801	AAACCTTTCG	GCCGTCTGGC	AGGAAATGGA	AATTATGAAT	CTGGTTTCCC
	851	TGCGTAAGCC	GGATGATGCC	TATGCGCGTT	TGCAAGTGCT	GTTGGAACAC
45	901	AACCCGAATG	CAAACTGTAT	TATTCAGGCG	CGCATATTGG	GCGCAAAACG
	951	AAAAGAAGGT	GCGTCCGTTA	TCGACGGCTA	CGCCGAAAAG	GCATACGGCA
	1001	GGGGGACGGG	GGAACAGCGG	GGCagggcgg	cAATgaacgc	GGCGATGATA
	1051	TATGCCGACC	GCAGGGATTA	CGCCAAAGTC	AGGCAGTGCT	TGAAAAAAGT
	1101	GTCCGCGCCG	GAATACCTGT	TCGACAAAGG	CGTGCTGGCG	GTCGCGCGCG
50	1151	CTGCCGAATT	GGACGGAGGC	CGGGCGGCTT	TGCGGCAGAT	CGGCAGGGTG
	1201	CGGAAACTTC	CCGAACAGCA	GGGGCGGTAT	TTTACGCGCA	ACAATTGTCT
	1251	CAAAATACAG	ATGCTCGCCC	TGTCGAAGCT	GCCCGACAAA	CGGGAAGCCC
	1301	TGATCGGGCT	GAACAACATC	ATCGCCAAAC	TTTCGGCGCG	GGGAAGCACG
	1351	GAACCTTTGG	CGGAAGCATT	GGCACAGCGT	TCCATTATTT	ACGaacAGTT
55	1401	cggCAAACGG	GGAAAAATGA	TTGCCGACCT	tgaAAcCgcg	CTCAAACTTA
	1451	CGCCCGATAA	TGCACAAATT	ATGAATAATC	TGGGCTACAG	CCTGCTTTCC
	1501	GATTCCAAAC	GTTTGAGACA	GGGTTTCGCC	CTGCTTCAGA	CGGCATACCA
	1551	AATCAACCCG	GACGATACCG	CCGTTAACGA	CAGCATAGGC	TGGGCGGTAT
	1601	ACCTGAAAGG	CGACGcggaA	AGCGCGCTGC	CGTATCTGcg	gtaattcggtt
	1651	aAAAACGACC	CCGAGCCCGA	AGTTGCCGCC	CATTTGGGGC	AAGTtTGTGTG

1701 GGCATTGGGC GAACGCGATC AGGCGGTTGA CGTATGGACG CAGGCGGCAC
 1751 ACCTTAGGGG AGACAAGAAA ATATGGCGGG AGACGCTCAA ACGCTACGGA
 1801 ATCGCCTTGC CCGAGCCTTC CCGAAAACCC CGGAAATAA

This encodes a protein having amino acid sequence <SEQ ID 48>:

5 1 MLPARFTILS VLAAALLAGQ AYAAGAADVE LPKEVGKVLK RHRRYSEEEI
 51 KNERARLAHV GERNRVFTL LGGETALQKG QAGTALATYM IMLERTKSPE
 101 VAERALEMAV SLNAFEQAEM IYQKWRQIEP IPGEAQKPAG WLRNVLKEGG
 151 NQHLDDLKEV LAQSDDDVQKR RIFLLLVQAA VQQGGVAQKA SKAVRRAALK
 10 201 YEHLPEAAVA DAVFGVQGRE KEKAIEALQR LAKLDTEILP PTLMTLRLTA
 251 RKYPEILDGF FEQTDTONLS AVWQEMEIMN LVSLRKPDDA YARLNVLLH
 301 NPNANLYIQ AILANRKEG ASVIDGYAEK AYGRGTGEOR GRAAMTAAMI
 351 YADRRDYAKV RQWLKVSAP EYLFDKGVLA AAAAAELDGG RAALRQIGRV
 401 RKLPEQQGRY FTADNLKIQ MLALSKLPDK REALIGLNNI IAKLSAAGST
 451 EPLAEALAQR SIIYEQFGKR GKMIADLETA LKLTPDNAQI MNNLGYSLLS
 15 501 DSKRLDEGFA LLQYAYQINP DDTAVNDSIG WAYYLGKDAE SALPYLRYSF
 551 ENDPEPEVAA HLGEVLWALG ERDQAVDVWT QAAHLRGDKK IWRETLKRYG
 601 IALPEPSRKP RK*

ORF9ng and ORF9-1 show 88.1% identity in 614 aa overlap:

20	orf9-1.pep	MLPNRKFMLTVLTATLIAGQVSAAGGGAGDMKQPKVEVGKVFRRKQRYSEEEIKNERARLA
	orf9ng-1	MLPARFTILSVLAAALLAGQAYAG--AADVELPKEVGKVLKRRYSEEEIKNERARLA
		10 20 30 40 50 60
25	orf9-1.pep	AVGERVNIQIFTLGGETALQKGQAGTALATYMLMLERTKSPEVAERALEMAVSLNAFEQA
	orf9ng-1	AVGERVNRVFTLLGGETALQKGQAGTALATYMLMLERTKSPEVAERALEMAVSLNAFEQA
		60 70 80 90 100 110
30	orf9-1.pep	EMIIYQKWRQIEPIPGKAQKRAGWLRNVLRERGNQHLGDEEVLAQADEGQNRVFLLLAQ
	orf9ng-1	EMIIYQKWRQIEPIPGEAQKPAGWLRNVLKEGGNQHLDGLKEVLAQSDDDVQKRIFLLLVQ
35		120 130 140 150 160 170
40	orf9-1.pep	AAVQQDGLAQKASKAVRRAALKYEHLPAAVADVFSVQGREKEKAIGALQRLAKLDTEI
	orf9ng-1	AAVQQGGVAQKASKAVRRAALKYEHLPAAVADAVFGVQGREKEKAIEALQRLAKLDTEI
		180 190 200 210 220 230
45	orf9-1.pep	LPPTLMTLRLTARKYPEILDGFFEQTDTQNL SAVWQEMEIMNLVSLHRLDDAYARLNVLL
	orf9ng-1	LPPTLMTLRLTARKYPEILDGFFEQTDTQNL SAVWQEMEIMNLVSLRKPDDAYARLNVLL
		240 250 260 270 280 290
50	orf9-1.pep	ERNPNADLYIQAILAANRKEGASVIDGYAEKAYGRGTGEORSRALTAAMMYADRRDYA
	orf9ng-1	EHNPNANLYIQAILAANRKEGASVIDGYAEKAYGRGTGEORGRAAMTAAMIYADRRDYA
		300 310 320 330 340 350
55	orf9-1.pep	KVRQWLKKSVAPEYLFDKGVLA AAAA AVELDGGRAALRQIGRVRLPEQQGRYFTADNL SK
	orf9ng-1	KVRQWLKKSVAPEYLFDKGVLA AAAAA AVELDGGRAALRQIGRVRLPEQQGRYFTADNL SK
		360 370 380 390 400 410
60	orf9-1.pep	IQMLALSKLPDKREALRGLDKIIEKPPAGSNTLQAEALVQSVVYDRLGKRRKMISDL E
	orf9ng-1	IQMLALSKLPDKREALIGLNNIIAKLSAAGSTEPLAEALAQRSIIYEQFGKRKGMIA DLE
65		420 430 440 450 460 470
		490 500 510 520 530 540

orf9-1.pep RAFRLAPDNAQIMNNLGYSLLTDSKRLDEGFALLQTAYQINPDDTAVNDSIGWAYYLKGD
 orf9ng-1 TALKLTPDNAQIMNNLGYSLLSDSKRLDEGFALLQTAYQINPDDTAVNDSIGWAYYLKGD
 480 490 500 510 520 530
 550 560 570 580 590 600
 orf9-1.pep AESALPYLRYSFENDEPEEVAHAHLGEVLWALGERDQAVDVWVQAHLRGDKKIWRKTLKR
 orf9ng-1 AESALPYLRYSFENDEPEEVAHAHLGEVLWALGERDQAVDVWVQAHLRGDKKIWRKTLKR
 540 550 560 570 580 590
 610
 orf9-1.pep HGIALPQPSRKPRKX
 orf9ng-1 YGIALPEPSRKPRKX
 600 610

In addition, ORF9ng shows significant homology with a hypothetical protein from *P.aeruginosa*:

sp|P42810|YHE3_PSEAE HYPOTHETICAL 64.8 KD PROTEIN IN HEMM-HEMA INTERGENIC REGION (ORF3)
 >gi|1072999|pir||S49376 hypothetical protein 3 - Pseudomonas aeruginosa >gi|557259 (X82071) orf3 [Pseudomonas aeruginosa] Length = 576
 Score = 128 bits (318), Expect = 1e-28
 Identities = 138/587 (23%), Positives = 228/587 (38%), Gaps = 125/587 (21%)
 Query: 67 VFTLLGGETALQKQAGTALATYMLMLERTKSPEVAERALEMAVSLNAFEQAEMIYQKWR 126
 +++LL E A Q+ + AL+ Y++ ++T+ P V+ERA +A L A ++A W
 Sbjct: 53 LYSLLVAELAGQRNRFDIASNYVVQAQKTRDPGVSEAFRIAIEYLGADQEAALDTSLLWA 112
 Query: 127 QIEPIPGEAQKPGAG-----WLRNVLKEGGNQHL DGLKEVLAQSDDVQKRR 172
 + P +AQ+ A ++ VL G+ H D L A++D + +
 Sbjct: 113 RSAPDNLDAQRAAIIQLARAGRYEESMYMEKVLNCGDTHDFLALSAAETDPDTRAGL 172
 Query: 173 FXXXXXXXXXXXXXXXXKASKAVRRAALKYEHLEAAVADAVFGVQGREKEKAIEALQRLA 232
 ++ KY + + A+ Q ++A+ L+ +
 Sbjct: 173 L-----QSFHLLKKYPNNGQLLFGKALLLQDGRPDALTLLEDNS 214
 Query: 233 KLDTEILPPTLMTLRLTARK-----YPEILDGFFEQTDTQNL SAVWQEMEIMNLVSLRKP 287
 E+ P L + L + K P + G E D + + + + LV +
 Sbjct: 215 ASRHEVAPLLLSRRLQSMKRSDEALPLLKAGIKEHPDDKRVRLAYARL----LVEQNRL 270
 Query: 288 DDAYARLNVLLLEHNP-----ANLYIQAAI----- 312
 DDA A L++ P+ A +Y++ +
 Sbjct: 271 DDAKAEFAGLVQFPDDDDDLRFLSALVCLAEQAQWDEARIYLEELVERDSHVDAAHFNLG 330
 Query: 313 -LAANRKEGASVIDGYAEKAYGRGTGEQGRGAAMTAAMIYADRRDYAKVRQWLKKVSAPE 371
 LA +K+ A +D YA+ G G + T ++ A R D A R + P+
 Sbjct: 331 RLAEQKDTARALDEYAQ--VGPGNDFLPAQLRQTDVLLKAGRVDEAAQRLDKARSEQPD 388
 Query: 372 YLFDKXXXXXXXXXXXXXXXXXQIGRVRKLPEQQGRYFTADNLSKIQLMLSKLPDKR 431
 Y A L I+ ALS +
 Sbjct: 389 Y-----AIQLYLIEAEALSNNNDQOE 408
 Query: 432 EALIGLNNIIAKLSAAGSTEPLAEALAQRSIIYEQFGKRGKMIADLETALKLTPDNAQIM 491
 +A + + + E L L RS++ E+ +M DL + PDNA +
 Sbjct: 409 KAWQAIQEGLKQYP----EDL-NLLYTRSMLEAKRNDLAQMEKDLRFVIAREPDMAMAL 462
 Query: 492 NNLGYSLLSDSKRLDEGFALLQTAYQINPDDTAVNDSIGWAYYLKGD AESALPYLRYSE 551
 N LGY+L + R E L+ A+++NPDD A+ DS+GW Y +G A YLR + +
 Sbjct: 463 NALGYTLADRTTRYGEARELILKAHKLNPDDPAILD SMGWINYRQGLADAERYLRQALQ 522
 Query: 552 NDPEPEVAHAHLGEVLWALGERDQAVDVWVQAHLRGDKKIWRKTLKR 598
 P+ EVAAHLGEVLWA G + A +W + + D + R T+KR
 Sbjct: 523 RYPDHEVAHAHLGEVLWAQGRQGDARAIWREYLDKQPDSDVLRRTIKR 569
 gi|2983399 (AE000710) hypothetical protein [Aquifex aeolicus] Length = 545
 Score = 81.5 bits (198), Expect = 1e-14
 Identities = 61/198 (30%), Positives = 98/198 (48%), Gaps = 19/198 (9%)
 Query: 408 GRYFTADNL-SKIQLMLSKLPDKREALIGLNNIIAKLSAAGSTEPLAEALAQ----- 459
 G Y A L K ++LA PDK+E L + +K + + L +

Sbjct: 335 GNYEDAKRLIEKAKVLA----PDKKEILFLEADYYSKTKQYDKALEILKKLEKDYPNDSR 390
 Query: 460 ----RSIIYEQFGKRGMIADELTALKLTPDNAQIMNNLGYSLLS--DSKRLDEGFALLQ 513
 +I+Y+ G L A++L P+N N LGYSLL +R++E L++
 5 Sbjct: 391 VYFMEAIVYDNLGDIKNAEKALRKAIELDPENPDYNYLGYSLLLWYGKERVEAEELIK 450
 Query: 514 TAYQINPDDTAVNDSIGWAYYLKGD AESALPYLRYSF-ENDPEPEVA AHLGEVLWALGER 572
 A + +P++ A DS+GW YYLKG D E A+ YL + E +P V H+G+VL +G +
 10 Sbjct: 451 KALEKDPENPAYIDSMGWVYYLKG DYERAMQYLLKALREAYDDFV VNEHVGDVLLKMGYK 510
 Query: 573 DQAVDVWTQAAHLRGDKK 590
 ++A + + +A L + K
 Sbjct: 511 EEARNYYERALKLLEEGK 528

- 15 Based on this analysis, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 7

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 49>:

20 1 AACCTCTACG CCGGCCCGCA GACCACATCC GTCATCGCAA ACATCGCCGA
 51 CAACCTGCAA CTGGCCAAAG ACTACGGCAA AGTACACTGG TTCGCCTCCC
 101 CGCTCTTCTG GCTCCTGAAC CAACTGCACA ACATCATCGG CAACTGGGGC
 151 TGGGCGATTA TCGTTTAAAC CATCATCGTC AAAGCCGTAC TGTATCCATT
 201 GACCAACGCC TCTTACCGCT CTATGGCGAA AATGCGTGCC GCCGCACCCA
 251 AACTGCAAGC CATCAAAGAG AAATACGGCG ACGACCGTAT GGCGCAACAA
 25 301 CAGGCGATGA TGCAGCTTTA CACAGACGAG AAAATCAACC CGaCTGGGCG
 351 GCTGCCTGCC TATGCTGTTG CAAATCCCCG TCTTCATCGG ATTGTATTGG
 401 GCATTGTTCG CCTCCGTAGA ATTGCGCCAG GCACCTTGCC TGGGTGGAT
 451 TACCGACCTC AGCCGCGCCG ACCCTACTA CATCCTGCCC ATCATTATGG
 501 CGGCAACGAT GTTCGCCCAA ACTTATCTGA ACCCGCCGCC GAaCGACCCG
 30 551 ATGCagGCGA AAATGATGAA AATCATGCCG TTGGTTTTCT CsGwCrTGT
 601 CTTCTTCTTC CCTGCCGgks TGGTATTGTA CTGGGTAGTC AACAACTCC
 651 TGACCATCGC CCAGCAATGG CACATCAACC GCAGCATCGA AAAACAACGC
 701 GCCCAAGGCG AAGTCGTTTC CTA

This corresponds to the amino acid sequence <SEQ ID 50; ORF11>:

35 1 ..NLYAGPQTS VIANIADNLQ LAKDYGKVHW FASPLFWLLN QLHNIIGNWG
 51 WAIIVLTIIV KAVLYPLTNA SYRSMAMRA AAPKLQAIKE KYGDDRMAQQ
 101 QAMMQLYTDE KINPLGGCLP MLLQIPVFIG LYWALFASVE LRQAPWLGI
 151 TDLRADPYY ILPIIMAATM FAQTYLNPPP TDPMQAKMMK IMPLVFSXXF
 201 FFFPAGXVLY WVVNNLLTIA QQWHINRSIE KQRAQGEVVS *

- 40 Further sequence analysis revealed the complete DNA sequence <SEQ ID 51>:

1 ATGGATTTTA AAAGACTCAC GCGGTTTTTC GCCATCGCGC TGGTGATTAT
 51 GATCGGCTGG GAAAAGATGT TCCCCACTCC GAAGCCAGTC CCCGCGCCCC
 101 AACAGGCAGC ACAACAACAG GCCGTAACCG CTTCCGCCGA AGCCGCGCTC
 151 GCGCCCGCAA CGCCGATTAC CGTAACGACC GACACGGTTC AAGCCGTCAT
 45 201 TGATGAAAAA AGCGCGGACC TGCGCGGCT GACCTGCTC AAATACAAAG
 251 CAACCGCGCA CGAAAATAAA CCGTTCATCC TGTTTGGCGA CGGCAAGAA
 301 TACACCTACG TCGCCCAATC CGAACTTTTG GACGCGCAGG GCAACAACAT
 351 TCTAAAGGC ATCGGCTTTA GCGCACCAGG AAAACAGTAC AGCTTGAAG
 401 GCGACAAAGT TGAAGTCCCG CTGAGCGCGC CTGAAACAGC CGGTCTGAA
 50 451 ATCGACAAAG TTTATACTTT CACCAAAGGC AGCTATCTGG TCAACGTCCG
 501 CTTGACATC GCCAACGGCA GCGGTCAAAC CGCCAACCTG AGCGCGGACT
 551 ACCGCATCGT CCGCGACCAC AGCGAACCCG AGGGTCAAGG TTACTTTACC
 601 CACTCTTACG TCGGCCCTGT TGT'TTATACC CCTGAAGGCA ACTTCCAAA
 651 AGTCAGCTTT TCCGACTTGG ACGACGATGC CAAATCCGGC AAATCCGAGG
 55 701 CCGAATACAT CCGCAAAACC CCGACCGGCT GGCTCGGCAT GATTGAACAC
 751 CACTTCATGT CCACCTGGAT TCTCCAACCT AAAGCGAGAC AAAGCGTTTG
 801 CGCCGCGAGC GAGTGAACA TCGACATCAA ACGCCGCAAC GACAAGCTGT
 851 ACAGCAACCAG CGTCAGCGTG CCTTTAGCCG CCATCCAAAA CGGCGCGAAA
 901 GCCGAAGCCT CCATCAACCT CTACGCCGGC CCGCAGACCA CATCCGTCAT
 60 951 CGCAACATC GCCGACAACC TGCAACTGGC CAAAGACTAC GGCAAGTAC

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1001 ACTGGTTCGC CTCCTCGCTC TTCTGGCTCC TGAACCAACT GCACAACATC
 1051 ATCGGCAACT GGGGCTGGGC GATTATCGTT TTAACCATCA TCGTCAAAGC
 1101 CGTACTGTAT CCATTGACCA ACGCCTCTTA CCGCTCTATG GCGAAAATGC
 1151 GTGCCGCCGC ACCCAAATG CAAGCCATCA AAGAGAAATA CGGCGACGAC
 1201 CGTATGGCGC AACAAACAGC GATGATGCAG CTTTACACAG ACGAGAAAAT
 1251 CAACCCGCTG GCGGGCTGCC TGCCTATGCT GTTGCAAATC CCCGTCTTCA
 1301 TCGGATTGTA TTGGGCATTG TTCGCCTCCG TAGAATTGCG CCAGGCACCT
 1351 TGGCTGGGTT GGATTACCGA CCTCAGCCGC GCCGACCCCT ACTACATCCT
 1401 CCCCATCATT ATGGCGGCAA CGATGTTCCG CCAAACCTTAT CTGAACCCGC
 1451 CGCCGACCGA CCCGATGCAG GCGAAAATGA TGAATCAT GCGGTTGGTT
 1501 TTCTCCGTC TGTCTTCTT CTTCCCTGCC GGTCTGGTAT TGTACTGGGT
 1551 AGTCAACAAC CTCCTGACCA TCGCCAGCA ATGGCACATC AACCGCAGCA
 1601 TCGAAAAACA ACGCGCCCAA GCGGAAGTCG TTTCCTAA

This corresponds to the amino acid sequence <SEQ ID 52; ORF11-1>:

15 1 MDFKRLTAFF AIALVIMIGW EKMFPKPKPV PAPQQAQQQ AVTASAEAL
 51 APATPITVTT DTVQAVIDEK SGDLRRLTLL KYKATGDENK PFILFGDGKE
 101 YTVVAQSELL DAQGNILKG IGFSAPKKQY SLEGDKVEVR LSAPETRGLK
 151 IDKVYTFKGY SYLVNVRFDI ANGSGQTANL SADYRIVRDH SEPEGQGYFT
 201 HSYVGPVYPT PEGNFQKVSF SLDLDDAKSG KSEAERYIKT PTGWLGMIEH
 251 HFMSTWILQP KGRQSVCAAG ECNIDIKRRN DKLYSTSVSV PLAAIQNGAK
 301 AEASINLYAG PQTTSVIANI ADNLQAKDY GKVHWFASPL FWLLNQLHNI
 351 IGNWGWAIIV LTIIVKAVLY PLTNASYRSM AKMRAAPKL QAIKEKYGDD
 401 RMAQQQAMMQ LYTDEKINPL GGCLPMLLQI PVFIGLYWAL FASVELRQAP
 451 WLGWITDLR ADPYYILPII MAATMFAQTY LNPPPTDPMQ AKMMKIMPLV
 25 501 FVMFFFFFA GLVLYWVNN LLTIAQQWHI NRSIEKQRAQ GEVVS*

Computer analysis of this amino acid sequence gave the following results:

Homology with a 60kDa inner-membrane protein (accession P25754) of *Pseudomonas putida*

ORF11 and the 60kDa protein show 58% aa identity in 229 aa overlap (BLASTp).

30 ORF11 2 LYAGPQTTSVIANIADNLQAKDYGVHWFASPLFWLLNQLHNIIGNWGWAIIVLTIIVK 61
 LYAGP+ S + ++ L+L DYG + + A P+FWLL +H+++GNWGW+IIVLT+++K
 60K 324 LYAGPKIQSKLKEKELSPGLELTVDYGFLLWFIAPFIWLLQHIHSLGNWGSIIIVLTMLIK 383
 ORF11 62 AVLYPLTNASYRSMAMRAAPKLQAIKEKYGDDRRXXXXXXXXXXLYTDEKINPLGGCLPM 121
 + +PL+ ASYRSMA+MRA APKL A+KE++GDDR LY EkinPLGGCLP+
 35 60K 384 GLFFPLSAASYRSMARMRAVAPKLAALKERFGDDRQKMSQAMMELYKKEKINPLGGCLPI 443
 ORF11 122 LLQIPVFIGLYWALFASVELRQAPWLGWITDLRADPYYILPII MAATMFAQTYLNPPPT 181
 L+Q+PVF+ LYW L SVE+RQAPW+ WITDLS DP++ILPIIM ATMF Q LNP P
 40 60K 444 LQMPVFLALYWVLLSEVMRQAPWILWITDLSIKDPFFILPIIMGATMFIQQRNPTPP 503
 ORF11 182 DPMQAKMMKIMPLVXXXXXXXXXVAGXVLYWVNNLLTIAQQWHINRSIE 230
 DPMQAK+MK+MP++ PAG VLYWVNN L+I+QW+I R IE
 60K 504 DPMQAKVMKMMPIIFTFFFLWFPAGLVLYWVNNCLSSISQQWYITRRIE 552

45 Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF11 shows 97.9% identity over a 240aa overlap with an ORF (ORF11a) from strain A of *N. meningitidis*:

50 orf11.pep 10 20 30
 NLYAGPQTTSVIANIADNLQAKDYGVHWF
 orf11a IKRRNDKLYSTSVSVPLAAIQNGAKSXASINLYAGPQTTSVIANIADNLQAKDYGVHWF
 280 290 300 310 320 330
 55 orf11.pep 40 50 60 70 80 90
 FASPLFWLLNQLHNIIGNWGWAIIVLTIIVKAVLYPLTNASYRSMAMRAAPKLQAIKE
 orf11a FASPLFWLLNQLHNIIGNWGWAIIVLTIIVKAVLYPLTNASYRSMAMRAAPKLQAIKE
 340 350 360 370 380 390

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		100	110	120	130	140	150
orf11.pep		KYGD	DRMA	QQQAM	QLYT	DEKIN	PLGGCLPMLLQIPVFIGLYWALFASVELRQAPWLGWI
5	orf11a	KYGD	DRMA	QQQAM	QLYT	DEKIN	PLGGCLPMLLQIPVFIGLYWALFASVELRQAPWLGWI
		400	410	420	430	440	450
		160	170	180	190	200	210
orf11.pep		TDLS	RADPY	YILPI	IMAAT	MFAQ	TYLNP
10	orf11a	TDLS	RADPY	YILPI	IMAAT	MFAQ	TYLNP
		460	470	480	490	500	510
		220	230	240			
orf11.pep		WVVN	NLLT	IAQQ	WHIN	RSIE	KQRAQGEVVSX
15	orf11a	WVIN	NLLT	IAQQ	WHIN	RSIE	KQRAQGEVVSX
		520	530	540			

The complete length ORF11a nucleotide sequence <SEQ ID 53> is:

	1	ANGG	ATTTTA	AAAG	ACTCAC	NGNG	TTTTTC	GCCAT	CGCAC	TGGT	GATTAT
20	51	GATC	GGATNG	NAAANG	ATGT	TCCCC	ACTCC	GAAG	CCCGTC	CCCG	CGCCCC
	101	AACAG	ACGGC	ACAACA	ACAG	GCCGT	AANCG	CTTCG	CCCGA	AGCC	GCGCTC
	151	GCGCC	CGNAN	CGCCG	ATTAC	CGTAA	CGACC	GACAC	GGTTC	AAGCC	GTCCAT
	201	TGAT	GAAAAA	AGCGG	CGACC	TGCGC	CGGCT	GACCT	GCTC	AAAT	ACAAAG
25	251	CAACC	GGCGA	CNAAA	ATAAA	CCGTT	CATCC	TGTTT	GGCGA	CGGCA	AAANAA
	301	TACAC	CTACN	TCGCC	CCANTC	CGAAC	TTTTG	GACGC	GCAGG	GCAAC	ACACAT
	351	TCTAA	AAGGC	ATCGG	CTTTA	GCGCA	CCGAA	AAAAC	AGTAC	AGCT	TGGAAG
	401	GCGAC	AAAGT	TGAAG	TCCGC	CTGAG	CGCAC	CTGAA	ACACG	CGGT	CTGAAA
	451	ATCGA	CAAAG	TTTAT	ACTTT	CACCA	AAGGC	AGCTA	TCTGG	TCAAC	GTCCG
30	501	CTTCG	ACATC	GCCAA	CGGCA	GCGGT	CAAAC	CGCCA	ACCTG	AGCG	CGGACT
	551	ACCGC	ATCGT	CCGCG	ACCAC	AGCGA	ACCCG	AGGGT	CAAGG	CTACT	TTTACC
	601	CACCT	CTTACG	TCGGC	CCCTGT	TGTTT	TATACC	CCTGA	AGGCA	ACTT	CCAAAA
	651	AGTC	AGCTTC	TCCGA	CTTGG	ACGAC	GATGC	CAANT	CCGGN	AAAT	CCGAGG
	701	CCGA	ATACAT	CCGCA	AAAACC	CNGAC	CGGCT	GGCTC	GGCAT	GATT	GAAACAC
35	751	CACCT	CATGT	CCACT	TGGAT	CCTCC	AACCC	AAAGG	CGGAC	AAAG	CGTTTTG
	801	CGCCG	CTGGC	GACTG	CCNGTA	TNGAC	ATCAA	ACGCG	CAAC	GACA	AGCTGT
	851	ACAGC	ACCAG	CGTC	AGCGTG	CCTTT	AGCCG	CTATC	CAAAA	CGGT	GCGAAA
	901	TCCNA	AGCCT	CCATC	AAACCT	CTACG	CCGGC	CCAC	AGACCA	CATC	NGTTAT
	951	CGCAA	ACATC	GCCGA	CAACC	TGCA	ACTGN	CAAAG	ACTAC	GGCA	AAAGTAC
40	1001	ACTGG	TTCGC	CTCCC	CCCTC	TTTTG	GCTTT	TGAAC	CAACT	GCAC	AACATC
	1051	ATCGG	CAACT	GGGGT	TGGGC	GATT	ATCGTT	TTAAC	CATCA	TCGT	CAAAGC
	1101	CGTAC	TGTAT	CCATT	TGACCA	ACGCT	CTCTTA	CCGTC	CGATG	GCGAA	AATGC
	1151	GTGCG	CCCGC	GCCAA	AACTG	CAAGC	CATCA	AAGAG	AAATA	CGGC	GACGAC
	1201	CGTAT	TGGCG	AGCAA	CAAGC	CATG	ATGCAG	CTTTA	CACAG	ACGAG	AAAAAT
45	1251	CAACC	CGCTG	GGCGG	CTGCC	TGCCT	ATGCT	GTTGC	AAATC	CCCG	TCTTCA
	1301	TCGG	ATTGTA	TTGGG	CATTG	TTCCG	CTCCG	TAGAA	TTGCG	CCAGG	CACCT
	1351	TGGCT	TGGGT	GGATT	ACCGA	CCTCA	GCCGC	GCCGA	CCNT	ACTAC	ATCCT
	1401	HSYV	GPVVYT	PEGN	FQKVSF	SDLD	DDAXSG	KSEAE	YIRKT	XTGW	LGMIEH
	1451	CGCC	GACCGA	CCCG	ATGCAG	GCGAA	AATGA	TGAAA	AATCAT	GCCT	TTGGTT
50	1501	NTNT	CNNNNA	NGTT	CTTCNN	CTTCC	CTGCC	GGCT	GCTGAT	TGTAC	TGGGT
	1551	GATCA	ACAAC	CTCCT	GACCA	TCGCC	CAGCA	ATGG	CACATC	AACCG	CAGCA
	1601	TCGAAA	AAACA	ACGCG	CCCAA	GGCGA	AGTCG	TTTCT	CTAA		

This encodes a protein having amino acid sequence <SEQ ID 54>:

	1	XDFK	RLTXFF	AIAL	VIMIGX	XXMF	PTPKPV	PAPQ	QTAQQQ	AVXA	SAEAAAL
55	51	APXX	PTVT	DTVQ	AVIDEK	SGDL	RRLTLL	KYKA	TGD	XNKP	PFILFGDGKX
	101	YTYX	AXSELL	DAQG	NNILKG	IGFS	APKKQY	SLEG	DKVEVR	LSAP	PETRGLK
	151	IDKV	YFTK	SYLV	NVRFDI	ANGS	GQTANL	SADY	RIVRDH	SEPE	GQGYFT
	201	HSYV	GPVVYT	PEGN	FQKVSF	SDLD	DDAXSG	KSEAE	YIRKT	XTGW	LGMIEH
	251	HFMST	WILQP	KGGQ	SVCAAG	DCXX	DIKRRN	DKLY	TSVSV	PLAA	IQNGAK
	301	SXAS	INLYAG	PQTT	SVIANI	ADNL	QLXKDY	GKVV	HFASPL	FWLL	NQLHNI
60	351	IGNW	GWAIIV	LTII	VKAVLY	PLTN	ASYRSM	AKMR	AAAPKL	QAIKE	KYGGD
	401	RMAQ	QQAMMQ	LYT	DEKINPL	GGCL	PMLLQI	PVFI	GLYWAL	FASV	ELRQAP
	451	WLGW	ITDLR	ADPY	YILPII	MAAT	MFAQTY	LNP	PP	TDPMQ	AKMMKIMPLV
	501	XSXX	FFXFFA	GLVL	YVWINN	LLTIA	QQWHI	NRSIE	KQRAQ	GEVVS*	

ORF11a and ORF11-1 show 95.2% identity in 544 aa overlap:

65

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5	orf11a.pep	XDFKRLTXFFAIALVIMIGXXXMFPTPKVPVPAQQTAAQQQAVXASAEALAPXXPITVTT
	orf11-1	MDFKRLTAFFAIALVIMIGWEKMFPTPKVPVPAQQAQQQAVTASAEALAPATPITVTT 10 20 30 40 50 60
10	orf11a.pep	DTVQAVIDEKSGDLRRLTLLKYKATGDXNKPFILFGDGKXYTYXASELLDAQGNNILKG
	orf11-1	DTVQAVIDEKSGDLRRLTLLKYKATGDENKPFILFGDGKEYTYVAQSELLDAQGNNILKG 70 80 90 100 110 120
15	orf11a.pep	IGFSAPKKQYSLEGDKVEVRLSAPETRGLKIDKVYTFTKGSYLVNVRFDIANGSGQTANL
	orf11-1	IGFSAPKKQYSLEGDKVEVRLSAPETRGLKIDKVYTFTKGSYLVNVRFDIANGSGQTANL 130 140 150 160 170 180
20	orf11a.pep	SADYRIVRDHSEPEGQGYFTHSYVGPVYTPEGNFQKVSFSDLDDDAKSGKSEAEYIRKT
	orf11-1	SADYRIVRDHSEPEGQGYFTHSYVGPVYTPEGNFQKVSFSDLDDDAKSGKSEAEYIRKT 190 200 210 220 230 240
25	orf11a.pep	XTGWLGMIEHHFMSTWILQPKGGQSVCAAGDCXXDIKRRNDKLYSTSVSVPLAAIQNGAK
	orf11-1	PTGWLGMIEHHFMSTWILQPKGRQSVCAAGECNIDIKRRNDKLYSTSVSVPLAAIQNGAK 250 260 270 280 290 300
30	orf11a.pep	SXASINLYAGPQTTSVIANIADNLQLKDYGKVHWFASPLFWLLNQLHNIIGNWGWAIIV :
	orf11-1	AEASINLYAGPQTTSVIANIADNLQLAKDYGKVHWFASPLFWLLNQLHNIIGNWGWAIIV 310 320 330 340 350 360
35	orf11a.pep	LTIIVKAVLYPLTNASYRSMAMRAAAPKLQAIKEKYGDDRMAMQAMQLYTDEKINPL
	orf11-1	LTIIVKAVLYPLTNASYRSMAMRAAAPKLQAIKEKYGDDRMAMQAMQLYTDEKINPL 370 380 390 400 410 420
40	orf11a.pep	GGCLPMLLQIPVFIGLYWALFASVELRQAPWLGWITDLSRADPYYILPIIIMAAATMFAQTY
	orf11-1	GGCLPMLLQIPVFIGLYWALFASVELRQAPWLGWITDLSRADPYYILPIIIMAAATMFAQTY 430 440 450 460 470 480
45	orf11a.pep	LNPPPTDPMQAKMMKIMPLVXSXXFFXFPAGLVLYWVNNLLTIAQQWHINRSIEKQRAQ
	orf11-1	LNPPPTDPMQAKMMKIMPLVFSVMFFFFPAGLVLYWVNNLLTIAQQWHINRSIEKQRAQ 490 500 510 520 530 540
50	orf11a.pep	GEVVSX
	orf11-1	GEVVSX

60 Homology with a predicted ORF from *N.gonorrhoeae*

ORF11 shows 96.3% identity over a 240aa overlap with a predicted ORF (ORF11.ng) from *N. gonorrhoeae*:

65	Orf11	NLYAGPQTTSVIANIADNLQLAKDYGKVHWFASPLFWLLNQLHNIIGNWGWAIIVLT 	57
	orf11ng	MAVNLYAGPQTTSVIANIADNLQLAKDYGKVHWFASPLFWLLNQLHNIIGNWGWAIIVLT 	60

	orf11	IIIVKAVLYPLTNASYRSMKMRRAAPKLQAIKEKYGDDRMAQQQAMMQLYTDEKINPLGG	117
	orf11ng	IIIVKAVLYPLTNASYRSMKMRRAAPELQTIKEKYGDDRMAQQQAMMQLFEDEEINPLGG	120
5	orf11	CLPMLLQIPVFIGLYWALFASVELRQAPWLGWITDLRADPYYILPIIMAATMFAQTYLN	177
	orf11ng	CLPMLLQIPVFIGLYWALFASVELRQAPWLGWITDLRADPYYILPIIMAATMFAQTYLN	180
10	orf11	PPPTDPMQAKMMKIMPLVFSXXFFFFPAGXVLYWVNNLLTIAQQWHINRSIEKQRAQGE	237
	orf11ng	PPPTDPMQAKMMKIMPLVFSVMFFFFPAGLVLYWVNNLLTIAQQWHINRSIEKQRAQGE	240
	orf11	VVS 240	
15	orf11ng	VVS 243	

An ORF11ng nucleotide sequence <SEQ ID 55> was predicted to encode a protein having amino acid sequence <SEQ ID 56>:

	1	MAVNLYAGPQ	TTSVIANIAD	NLQLAKDYGK	VHWFASPLFW	LLNQLHNIIG
	51	NWGWAIIVLT	IIIVKAVLYPL	TNASYRSMK	MRAAAPELQT	IKEKYGDDRM
20	101	AQQQAMMQLF	EDEEINPLGG	CLPMLLOIPV	FIGLYWALFA	SVELRQAPWL
	151	GWITDLRAD	PYYILPIIMA	ATMFAQTYLN	PPPTDPMQAK	MMKIMPLVFS
	201	VMFFFFPAGL	VLYWVNNLL	TIAQQWHINR	SIEKQRAQGE	VVS*

Further sequence analysis revealed the complete gonococcal DNA sequence <SEQ ID 57> to be:

	1	ATGGATTTTA	AAAGACTCAC	GGCGTTTTTC	GCCATCGCGC	TGGTGATTAT
25	51	GATCGGCTGG	GAATAAATGT	TCCCCACCCC	GAAACCCGTC	CCCGCGCCCC
	101	AACAGGCGGC	ACAAAAACAG	GCAGCAACCG	CTTCCGCCGA	AGCCGCGCTC
	151	GCGCCCGCAA	CGCCGATTAC	CGTAACGACC	GACACGGTTC	AAGCCGTTAT
	201	TGATGAAAAA	AGTGGCGACC	TGCGCCGGCT	GACCCTGCTC	AAATACAAAG
	251	CAACCGGCGA	CGAAACAAA	CCGTTCGTCC	TGTTTGGCGA	CGGCAAAGAA
30	301	TACACCTACG	TCGCCCAATC	CGAACTTTTG	GACGCGCAGG	GCAACAACAT
	351	TCTGAAAGGC	ATCGGCTTTA	GCGCACCAGG	AAAACAGTAC	ACCCTCAACG
	401	GCGACACAGT	CGAAGTCCGC	CTGAGCGCGC	CCGAAACCAA	CGGACTGAAA
	451	ATCGACAAAG	TCTATACCTT	TACCAAAGAC	AGCTATCTGG	TCAACGTCCG
	501	CTTCGACATC	GCCAACGGCA	GCGGTCAAAC	CGCCAACCTG	AGCGCGGACT
35	551	ACCGCATCGT	CCGCGACCAC	AGCGAACCCG	AGGGTCAAGG	CTACTTTACC
	601	CACCTTTACG	TCGGCCCTGT	TGTTTATACC	CCTGAAGGCA	ACTTCCAAAA
	651	AGTCAGCTTC	TCCgacTTgg	acgACGATGC	gaaaTccggc	aaATccgagg
	701	ccgaatacat	CCGCAAAACC	ccgaccgggt	ggctcggcat	gattgaacac
	751	cacttcatgt	ccacttggat	cctccAAcct	aaaggcggcc	aaaaagtttg
40	801	cgcccaggga	gactgcccga	tcgacattaa	aCgcccgaac	gacaagctgt
	851	acagcgcaag	cgtcagcgtg	cctttaaccg	ctatcccaac	ccggggggcca
	901	aaaccgaaaa	tgggcgTCAA	CCTGTATGCC	GGTCCGCAA	CCACATCCGT
	951	TATCGCAAAC	ATCGCcgacA	ACCTGCAACT	GGCAAAAGAC	TACGGTAAAG
	1001	TACACTGGTT	CGCATCGCCG	CTCTTCTGGC	TCCTGAACCA	ACTGCACAAC
45	1051	ATTATCGGCA	ACTGGGGCTG	GGCAATCGTC	GTTTGTGACCA	TCATCGTCAA
	1101	AGCCGTACTG	TATCCATTGA	CCAACGcttc	ctACCGTTTC	ATGGCGAAAA
	1151	TGCGTGccgc	cgcacCcaaA	CTGCAGACCA	TCAAAGAAAA	ATAcgGCGAC
	1201	GACCGTATGG	CGCAACAGCA	AGCGATGATG	CAGCTTTACA	AAGacgAGAA
	1251	AATCAACCCG	CTGGGCGGCT	GTctgcctat	gctgttgCAA	ATCCCCGTCT
50	1301	TCATCGGCTT	GTA CTGGGCA	TTGTTCGCCT	CCGTAGAATT	GCGCCAGGCA
	1351	CCTTGGCTGG	GCTGGATTAC	CGACCTCAGC	CGCGCCGACC	CCTACTACAT
	1401	CCTGCCCATC	ATTATGGCGG	CAACGATGTT	CGCCCAAACC	TATCTGAACC
	1451	CGCCGCGGAC	CGACCCGATG	CAGGCGAAAA	TGATGAAAAT	CATGCCGTTG
	1501	GTTTTCTCCG	TCATGTTCTT	CTTCTTCCCT	GCCGTTTGG	TTCTCTACTG
55	1551	GGTGGTCAAC	AACCTCCTGA	CCATCGCCCA	GCAGTGGCAC	ATCAACCGCA
	1601	GCATCGAAAA	ACAACGCGCC	CAAGCGGAAG	TCGTTTCCTA	A

This encodes a protein having amino acid sequence <SEQ ID 58; ORF11ng-1>:

	1	MDFKRLTAFF	AIALVIMIGW	EKMFPPTPKPV	PAPOQAAQKQ	AATASAEAL
60	51	APATPITVTT	DTVQAVIDEK	SGDLRRLTLL	KYKATGDENK	PFVLFQDGKE
	101	YTYVAQSELL	DAQGNNILKG	IGFSAPKKQY	TLNGDTVEVR	LSAPETNGLK
	151	IDKVYTFTKD	SYLVNVRFDI	ANGSGQTANL	SADYRIVRDH	SEPEQGQYFT
	201	HSYVGPVYYT	PEGNFQKVSF	SDLDDAKSG	KSEABYIRKT	PTGWLGMIEH
	251	HFMSTWILQP	KGGQNVCAQG	DCRIDIKRRN	DKLYSASVS	PLTAIPTRGP
	301	KPKMAVNLYA	GPQTTSVIAN	IADNLQLAKD	YGVHWFASP	LFWLLNQLHN

-90-

```

351 IIGNWGWAIV VLTIIIVKAVL YPLTNASYRS MAKMRAAAPK LQTIKEYGKD
401 DRMAQQQAMM QLYKDEKINP LGGCLPMLLO IPVFIGLYWA LFASVELRQA
451 PWLGWITDLS RADPYYILPI IMAATMFAQT YLNPPPTDPM QAKMMKIMPL
501 VFSVMFFFFFF AGLVLYWVWN NLLTIAQQWH INRSIEKQRA QGEVVS*

```

5 ORF11ng-1 and ORF11-1 shown 95.1% identity in 546 aa overlap:

```

10 orf11ng-1.pep      10      20      30      40      50      60
    MDFKRLTAFFAIALVIMIGWEKMFPTPKPVPAPQQAQQAATASAEALAPATPITVTT
    |||||
orf11-1      MDFKRLTAFFAIALVIMIGWEKMFPTPKPVPAPQQAQQAATASAEALAPATPITVTT
    10      20      30      40      50      60

15 orf11ng-1.pep      70      80      90     100     110     120
    DTVQAVIDEKSGDLRRLTLLKYKATGDENKPFVLFQDGKEYTYVAQSELLDAQGNNILKG
    |||||
orf11-1      DTVQAVIDEKSGDLRRLTLLKYKATGDENKPFVLFQDGKEYTYVAQSELLDAQGNNILKG
    70      80      90     100     110     120

20 orf11ng-1.pep      130     140     150     160     170     180
    IGFSAPKKQYTLNGDTVEVRLSAPETNGLKIDKVYFTKDSYLVNVRFDIANGSGQTANL
    |||||
orf11-1      IGFSAPKKQYTLNGDTVEVRLSAPETNGLKIDKVYFTKDSYLVNVRFDIANGSGQTANL
    130     140     150     160     170     180

25 orf11ng-1.pep      190     200     210     220     230     240
    SADYRIVRDHSEPEGQGYFTHSYVGPVYVTPEGNFQKVSFSDLDLDDAKSGKSEAEYIRKT
    |||||
orf11-1      SADYRIVRDHSEPEGQGYFTHSYVGPVYVTPEGNFQKVSFSDLDLDDAKSGKSEAEYIRKT
    190     200     210     220     230     240

30 orf11ng-1.pep      250     260     270     280     290     300
    PTGWLGMIEHHFMSTWILQPKGGQNVCAQGGCRIDIKRRNDKLYSASVSVPLTAIPTRGP
    |||||
orf11-1      PTGWLGMIEHHFMSTWILQPKGRQSVCAAGECNIDIKRRNDKLYSTSVSVPLAAIQN-GA
    250     260     270     280     290

35 orf11ng-1.pep      310     320     330     340     350     360
    KPKMAVNLYAGPQTTSVIANIADNLQAKDYGKVHWFASPLFWLLNQLHNIIGNWGWAIV
    | : : |||||
orf11-1      KAEASINLYAGPQTTSVIANIADNLQAKDYGKVHWFASPLFWLLNQLHNIIGNGWAI
    300     310     320     330     340     350

40 orf11ng-1.pep      370     380     390     400     410     420
    VLTIIIVKAVLYPLTNASYRSMAKMRAAAPKLQTIKEYGDDRMAQQQAMMQLYKDEKINP
    |||||
orf11-1      VLTIIIVKAVLYPLTNASYRSMAKMRAAAPKLQAIKEYGDDRMAQQQAMMQLYTDEKINP
    360     370     380     390     400     410

45 orf11ng-1.pep      430     440     450     460     470     480
    LGGCLPMLLOIPVFIGLYWALFASVELRQAPWLGWITDLSRADPYYILPI IMAATMFAQT
    |||||
orf11-1      LGGCLPMLLOIPVFIGLYWALFASVELRQAPWLGWITDLSRADPYYILPI IMAATMFAQT
    420     430     440     450     460     470

50 orf11ng-1.pep      490     500     510     520     530     540
    YLNPPPTDPMQAKMMKIMPLVFSVMFFFFFFPAGLVLYWVWNLLTIAQQWHINRSIEKQRA
    |||||
orf11-1      YLNPPPTDPMQAKMMKIMPLVFSVMFFFFFFPAGLVLYWVWNLLTIAQQWHINRSIEKQRA
    480     490     500     510     520     530

60 orf11ng-1.pep      QGEVVSX
    |||||
orf11-1      QGEVVSX
    540

```

65 In addition, ORF11ng-1 shows significant homology with an inner-membrane protein from the database (accession number p25754):

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```

              510      520      530      540
orf11ng-1.pep  SVMFFFFPAGLVLYWVVNNLLTIAQQWHINRSIEKQRAQGEVVSX
                : ::::| | | | | | | | | | | | | | | | | | | |
p25754         TFFFLWFEPAGLVLYWVVNNCLSSISQQWYITRRIEAATKKA
              520      530      540      550      560

```

Based on this analysis, including the homology to an inner-membrane protein from *P. putida* and the predicted transmembrane domains (seen in both the meningococcal and gonococcal proteins), it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

5 Example 8

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 59>:

```

1  ..GCCGTCTTAA TCATCGAATT ATTGACGGGA ACGGTTTATC TTTTGGTTGT
51  NAGCGCGGCT TTGGCGGGTT CGGGCATTGC TTACGGGCTG ACCGGCAGTA
101 CGCCTGCCGC CGTCTTGACC GNCGCTCTGC TTTCCGCGCT GGGTATTTNG
151 TTCGTACACG CCAAACCCGC CGTTAGAAAA GTTGAAACGG ATTCATATCA
201 GGATTGGAT GCCGACAAT ATGTCGAAAT CCTCCGNAC ACAGCGGGCA
251 ACCGTTACGA AGTT.TTTAT CGCGGTACG. ACTGGCAGGC TCAAAATACG
301 GGGCAAGAAG AGCTTGAACC AGGAACTCGC GCCCTCATTG TCCGCAAGGA
351 AGGCAACCTT CTTATTATCA CACACCCTTA A

```

15 This corresponds to the amino acid sequence <SEQ ID 60; ORF13>:

```

1  ..AVLIIELLTG TVYLLVVSAA LAGSGIAYGL TGSTPAAVLT XALLSALGIX
51  FVHAKTAVRK VETDSYQDL D AGQYVEILRH TGGNRYEVXY RGTWQAQNT
101 GQEELEPGTR ALIVRKEGNL LIITHP*

```

Further sequence analysis elaborated the DNA sequence slightly <SEQ ID 61>:

```

20 1  ..GCCGTCTTAA TCATCGAATT ATTGACGGGA ACGGTTTATC TTTTGGTTGT
51  nAGCGCGGCT TTGGCGGGTT CGGGCATTGC TTACGGGCTG ACCGGCAGTA
101 CGCCTGCCGC CGTCTTGACC GnCGCTCTGC TTTCCGCGCT GGGTATTTnG
151 TTCGTACACG CCAAACCCGC CGTTAGAAAA GTTGAAACGG ATTCATATCA
201 GGATTGGAT GCCGACAAT ATGTCGAAAT CCTCCGACAC ACAGCGGGCA
25 251 ACCGTTACGA AGTTTTtTAT CGCGGTACGc ACTGGCAGGC TCAAAATACG
301 GGGCAAGAAG AGCTTGAACC AGGAACTCGC GCCCTCATTG TCCGCAAGGA
351 AGGCAACCTT CTTATTATCA CACACCCTTA A

```

This corresponds to the amino acid sequence <SEQ ID 62; ORF13-1>:

```

30 1  ..AVLIIELLTG TVYLLVVSAA LAGSGIAYGL TGSTPAAVLT XALLSALGIX
51  FVHAKTAVRK VETDSYQDL D AGQYVEILRH TGGNRYEVFY RGTWQAQNT
101 GQEELEPGTR ALIVRKEGNL LIITHP*

```

Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF13 shows 92.9% identity over a 126aa overlap with an ORF (ORF13a) from strain A of *N.*

35 *meningitidis*:

```

                                     10      20      30      40      50
orf13.pep      AVLIIELLTGTVYLLVVSAA LAGSGIAYGLTGSTPAAVLT XALLSALGIXF
               |||
orf13a         MTVWFVA AVAVLIIELLTGTVYLLVVSAA LAGSGIAYGLTGSTPAAVLT AALLSALGIWF
               10      20      30      40      50      60

               60      70      80      90      100     110
orf13.pep      VHAKTAVRK VETDSYQDL DAGQYVEILRH TGGNRYEVXYRGTWQAQNT GQEELEPGTRA
               |||
orf13a         VHAKTAVGK VETDSYQDL DAGQYAEILRH AGGNRYEVFYRGTWQAQNT GQEELEPGTRA
               70      80      90      100     110     120

               120
orf13.pep      LIVRKEGNLLIITHPX
               |||

```

orf13a LIVRKEGNLLIIAKPX
130

The complete length ORF13a nucleotide sequence <SEQ ID 63> is:

```

5      1  ATGACTGTAT GGTGTGTTGC CGCTGTTGCC GTCTTAATCA TCGAATTATT
      51  GACGGGAACG GTTTATCTTT TGGTTGTCAG CGCGGCTTTG GCGGGTTCGG
     101  GCATTGCTTA CCGGCTGACC GGCAGCACGC CTGCCGCCGT CTTGACCGCC
     151  GCTCTGCTTT CCGCGCTGGG TATTTGGTTC GTACACGCCA AAACCGCCGT
     201  GGGAAAAGTT GAAACGGATT CATATCAGGA TTTGGATGCC GGGCAATATG
     251  CCGAAATCCT CCGGCACGCA GCGGCAACC GTTACGAAGT TTTTATCGC
    10  301  GGTACGCACT GGCAGGCTCA AAATACGGGG CAAGAAGAGC TTGAACCAGG
     351  AACGCGCGCC CTAATCGTCC GCAAGGAAGG CAACCTTCTT ATCATCGCAA
     401  AACCTTAA
  
```

This encodes a protein having amino acid sequence <SEQ ID 64>:

```

15      1  MTWVFVAAVA VLIIELLTGT VYLLVVSAA AGSGIAYGLT GSTPAAVLTA
      51  ALLSALGIWF VHAKTAVGKV ETDSYQDLDA QYAEILRHA GGNRYEVFYR
     101  GTHWQAQNTG QEELEPGTRA LIVRKEGNLL IIAKP*
  
```

ORF13a and ORF13-1 show 94.4% identity in 126 aa overlap

```

20      10      20      30      40      50      60
    orf13a.pep  MTWVFVAAVAVLIIELLTGTVYLLVVSAAALAGSGIAYGLTGSTPAAVLTAALLSALGIWF
    orf13-1      AVLIIELLTGTVYLLVVSAAALAGSGIAYGLTGSTPAAVLTXALLSALGIXF
                10      20      30      40      50

25      70      80      90      100     110     120
    orf13a.pep  VHAKTAVGKVETDSYQDLDAQYAEILRHAGGNRYEVFYRGTHWQAQNTGQEELEPGTRA
    orf13-1      VHAKTAVRKVETDSYQDLDAQYVEILRHTGGNRYEVFYRGTHWQAQNTGQEELEPGTRA
                60      70      80      90      100     110

30      130
    orf13a.pep  LIVRKEGNLLIIAKPX
    orf13-1      LIVRKEGNLLIIITHPX
                120
  
```

Homology with a predicted ORF from *N.gonorrhoeae*

ORF13 shows 89.7% identity over a 126aa overlap with a predicted ORF (ORF13.ng) from *N.gonorrhoeae*:

```

40      orf13      AVLIIELLTGTVYLLVVSAAALAGSGIAYGLTGSTPAAVLTXALLSALGIXF  51
    orf13ng  MTWVFVAAVAVLIIELLTGTVYLLVVSAAALAGSGIAYGLTGSTPAAVLTAALLSALGIWF  60

    orf13      VHAKTAVRKVETDSYQDLDAQYVEILRHTGGNRYEVXYRGTXWQAQNTGQEELEPGTRA  111
    orf13ng  VHAKTAVGKVETDSYQDLDTGKYAEILRYTGGNRYEVFYRGTHWQAQNTGQEVFEPGTRA  120

    orf13      LIVRKEGNLLIIITHP  126
    orf13ng  LIVRKEGNLLIIANP  135
  
```

The complete length ORF13ng nucleotide sequence <SEQ ID 65> is:

```

55      1  ATGACTGTAT GGTGTGTTGC CGCTGTTGCC GTCTTAATCA TCGAATTATT
      51  GACGGGAACG GTTTATCTTT TGGTTGTCAG CGCGGCTTTG GCGGGTTCGG
     101  GCATTGCCTA CCGGCTGACT GGCAGCACGC CTGCCGCCGT CTTGACCGCC
     151  GCACTGCTTT CCGCGCTGGG CATTTGGTTC GTACATGCCA AAACCGCCGT
     201  GGGAAAAGTT GAAACGGATT CATATCAGGA TTTGGATACC GGAAAATATG
     251  CCGAAATCCT CCGATACACA GCGGCAACC GTTACGAAGT TTTTATCGC
     301  GGTACGCACT GGCAGGCGCA AAATACGGGG CAGGAAGTGT TTGAACCGGG
     351  AACGCGCGCC CTCATCGTCC GCAAAGAAGG TAACCTTCTT ATCATCGCAA
     401  ACCCTTAA
  
```

This encodes a protein having amino acid sequence <SEQ ID 66>:

```

1  MTVWFVAAVA VLIIELLTGT VYLLVVSAAAL AGSGIAYGLT GSTPAAVLTA
51  ALLSALGIWF VHAKTAVGKV ETDYQDLDT GKYAEILRYT GGNRYEVFYR
101 GTHWQAQNTG QEVFEPGTRA LIVRKEGNLL IIANP*

```

5 ORF13ng shows 91.3% identity in 126 aa overlap with ORF13-1:

```

10 orf13-1.pep      10      20      30      40      50
      AVLIIELLTGTVYLLVVSAAALAGSGIAYGLTGSTPAAVLTXALLSALGIXF
orf13ng      MTVWFVAAVAVLIIELLTGTVYLLVVSAAALAGSGIAYGLTGSTPAAVLTAALLSALGIWF
      10      20      30      40      50      60

15 orf13-1.pep      60      70      80      90      100     110
      VHAKTAVRKVETDSYQDLDTAGQYVEILRHTGGNRYEVFYRGTHWQAQNTGQEELEPGTRA
orf13ng      VHAKTAVGKVETDSYQDLDTGKYAEILRYTGGNRYEVFYRGTHWQAQNTGQEVFEPGTRA
      70      80      90      100     110     120

20 orf13-1.pep      120
      LIVRKEGNLLIITHPX
orf13ng      LIVRKEGNLLIIANPX
      130

```

Based on this analysis, including the extensive leader sequence in this protein, it is predicted that ORF13 and ORF13ng are likely to be outer membrane proteins. It is thus predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 9

The following DNA sequence was identified in *N.meningitidis* <SEQ ID 67>:

```

30 1  ATGTWtGATT TCGGTTTrGG CGArCTGGTT TTTGTCGGCA TTATCGCCCT
51  GATWgtCCTC GGCCCCGAAC GCsTGCCCGA GGCCGCCCGC AyCGCCGGAC
101 GGcTCATCGG CAGGCTGCAA CGCTTTGTGCG GcAGCGTCAA ACAGGAATTT
151 GACACTCAAA TCGAACTGGA AGAACTGAGG AAGGCAAAGC AGGAATTGTA
35 201 AGCTGCCGcC GCTCAGGTTc GAGACAGCCT CAAAGAAACC GGTACGGATA
251 TGGAAGGCAA TCTGCACGAC ATTTCCGACG GTCTGAAGCC TTGGGAAAAA
301 CTGCCCGAAC AGCGGACACC TGCCGATTTC GGTGTCGATG AAAACGGCAA
351 TCCGCT.TCC CGATGCGGCA AACACCCATAT CAGACGGCAT TTCCGACGTT
401 ATGCCGTC..

```

This corresponds to the amino acid sequence <SEQ ID 68; ORF2>:

```

40 1  MXDFGLGELV FVGIIALIVL GPERXPEAAR XAGRLIGRLQ RFVGSVKQEF
51  DTQIELEELR KAKQEFEEAAA AQVRDSLKET GTDMEGNLHD ISDGLKPWEK
101 LPEQRTPADF GVDENGNPXS RCGKHPIRRH FFRYAV..

```

Further work revealed the complete nucleotide sequence <SEQ ID 69>:

```

45 1  ATGTTTGATT TCGGTTTGGG CGAGCTGGTT TTTGTCGGCA TTATCGCCCT
51  GATTGTCTCT GGCCCCGAAC GCCTGCCCGA GGCCGCCCGC ACCGCCGGAC
101 GGCTCATCGG CAGGCTGCAA CGCTTTGTGCG GCAGCGTCAA ACAGGAATTT
151 GACACTCAAA TCGAACTGGA AGAACTGAGG AAGGCAAAGC AGGAATTGTA
201 AGCTGCCGCC GCTCAGGTTc GAGACAGCCT CAAAGAAACC GGTACGGATA
50 251 TGGAAGGCAA TCTGCACGAC ATTTCCGACG GTCTGAAGCC TTGGGAAAAA
301 CTGCCCGAAC AGCGGACACC TGCCGATTTC GGTGTCGATG AAAACGGCAA
351 TCCGCTTCCC GATGCGGCAA ACACCCATAT AGACGGCATT TCCGACGTTA
401 TGCCGTCGCA ACGTTCCTAC GCTTCCGCCG AAACCCCTTG GGACAGCGGG
451 CAAACCGGCA GTACAGCCGA ACCCGCGGAA ACCGACCAAG ACCGCGCATG
501 GCGGGAATAC CTGACTGCTT CTGCCGCCGC ACCCGTCGTA CAGACCGTCG

```

551 AAGTCAGCTA TATCGATACT GCTGTTGAAA CGCCTGTTCC GCACACCACT
 601 TCCCTGCGCA AACAGGCAAT AAGCCGCAAA CGCGATTTTC GTCCGAAACA
 651 CCGCGCCAAA CCTAAATTGC GCGTCCGTAA ATCATAA

This corresponds to the amino acid sequence <SEQ ID 70; ORF2-1>:

5 1 MFDFGLGELV FVGIIALIVL GPERLPEAAR TAGRLIGRLQ RFVGSVKQEF
 51 DTQIELEELR KAKQEFEEAAA AQVRDSLKET GTDMEGNLHD ISDGLKPWEK
 101 LPEQRTPADF GVDENGNPFP DAANTLSDGI SDVMPSESY ASAETLGDSG
 151 QTGSTAEPAE TDQDRAWREY LTASAAAPVV QTVEVSYIDT AVETPVPHTT
 201 SLRKQAI SRK RDLRPKSRAK PKLRVRKS*

10 Further work identified the corresponding gene in strain A of *N.meningitidis* <SEQ ID 71 >:

1 ATGTTTGATT TCGGTTTGGG CGAGCTGGTT TTTGTCGGCA TTATCGCCCT
 51 GATTGTCTCT GGGCCCGAAC GCCTGCCCGA GGCCGCCCGC ACCGCCGGAC
 101 GGCTCATCGG CAGGCTGCAA CGCTTTGTCG GCAGCGTCAA ACAGGAATTT
 151 GACACGCAAA TCGAACTGGA AGAACTAAGG AAGGCAAAGC AGGAATTTGA
 15 AGCTGCCGCT GCTCAGGTTT GAGACAGCCT CAAAGAAACC GGTACGGATA
 251 TGGAGGGTAA TCTGCACGAC ATTTCCGACG GTCTGAAGCC TTGGGAAAAA
 301 CTGCCCGAAC AGCGCACGCC TGCTGATTTT GGTGTCGATG AAAACGGCAA
 351 TCCCTTTCCC GATGCGGCAA ACACCCATT AGACGGCATT TCCGACGTTA
 401 TGCCGTCCGA ACGTTCCTAC GCTTCCGCGG AAACCCTTGG GGACAGCGGG
 20 451 CAAACCGGCA GTACAGCCGA ACCCGCGGAA ACCGACCAAG ACCGTGCATG
 501 GCGGGAATAC CTGACTGCTT CTGCGCGCGC ACCGTCGTA CAGACCGTCG
 551 AAGTCAGCTA TATCGATACT GCTGTTGAAA CCCCTGTTCC GCATACCACT
 601 TCGCTGCGTA AACAGGCAAT AAGCCGCAAA CGCGATTTGC GTCCTAAATC
 651 CCGCGCCAAA CCTAAATTGC GCGTCCGTAA ATCATAA

25 This encodes a protein having amino acid sequence <SEQ ID 72; ORF2a>:

1 MFDFGLGELV FVGIIALIVL GPERLPEAAR TAGRLIGRLQ RFVGSVKQEF
 51 DTQIELEELR KAKQEFEEAAA AQVRDSLKET GTDMEGNLHD ISDGLKPWEK
 101 LPEQRTPADF GVDENGNPFP DAANTLLDGI SDVMPSESY ASAETLGDSG
 151 QTGSTAEPAE TDQDRAWREY LTASAAAPVV QTVEVSYIDT AVETPVPHTT
 30 201 SLRKQAI SRK RDLRPKSRAK PKLRVRKS*

The originally-identified partial strain B sequence (ORF2) shows 97.5% identity over a 118aa overlap with ORF2a:

		10	20	30	40	50	60
35	orf2.pep	MXDFGLGELVFVGIIALIVLGPXRXPEARXAGRLIGRLQRFVGSVKQEFDTQIELEELR					
	orf2a	MFDFGLGELVFVGIIALIVLGPRLPEAARTAGRLIGRLQRFVGSVKQEFDTQIELEELR					
		10	20	30	40	50	60
40	orf2.pep	KAKQEFEEAAAQVRDSLKETGTDMEGNLHDISDGLKPWEKLPEQRTPADFGVDENGNPXS					
	orf2a	KAKQEFEEAAAQVRDSLKETGTDMEGNLHDISDGLKPWEKLPEQRTPADFGVDENGNPFP					
		70	80	90	100	110	120
45	orf2.pep	RCGKHPIRRHFRRYAV					
	orf2a	DAANTLLDGISDVMPSESYASAETLGDSGQTGSTAEPAE TDQDRAWREYLTASAAAPVV					
		130	140	150	160	170	180

50 The complete strain B sequence (ORF2-1) and ORF2a show 98.2% identity in 228 aa overlap:

	orf2a.pep	MFDFGLGELVFVGIIALIVLGPRLPEAARTAGRLIGRLQRFVGSVKQEFDTQIELEELR	60
	orf2-1	MFDFGLGELVFVGIIALIVLGPRLPEAARTAGRLIGRLQRFVGSVKQEFDTQIELEELR	60
55	orf2a.pep	KAKQEFEEAAAQVRDSLKETGTDMEGNLHDISDGLKPWEKLPEQRTPADFGVDENGNPFP	120
	orf2-1	KAKQEFEEAAAQVRDSLKETGTDMEGNLHDISDGLKPWEKLPEQRTPADFGVDENGNPFP	120
60	orf2a.pep	DAANTLLDGISDVMPSESYASAETLGDSGQTGSTAEPAE TDQDRAWREYLTASAAAPVV	180

10

1	MFDFGLGELI	<u>FVGIIALIVL</u>	GPERLPEAAR	TAGRLIGRLQ	RFVGSVKQEL
51	DTQIELEELR	KVKQAFEEAA	AQVRDSLKET	DTDMQNSLHD	ISDGLKPWEK
101	LPEORTPADF	GVDEKGNLSL	RYGKHRIRRH	FRRYAV*	

	1	ATGTTTGATT	TCGGTTTGGG	CGAGCTGATT	TTTGTGCGCA	TTATCGCCCT
15	51	GATTGTCTT	GGTCCAGAAC	GCCTGCCCGA	AGCCGCCCGC	ACTGCCGGAC
	101	GGCTTTATCG	CAGGCTGCAA	CGCTTTGTAG	GAAGCGTCAA	ACAAGAACTT
	151	GACACTCAAA	TCCGAATGGA	AGAGCTGAGG	AAGGTCATGC	AGGCATTCGA
	201	AGCTGCCGCC	GCTCAGGTTT	GAGACAGCCT	CAAAGAAACC	GATACGGATA
20	251	TGCAGAACAG	TCTGCACGAC	ATTTCCGACG	GTCTGAAGCC	TTGGGAAAAA
	301	CTGCCCGAAC	AGCGCACGCT	tgccgatttc	gtGTCTGCATg	AAAaccggcaa
	351	tcccttctcc	gATACGGCAA	ACACCGTATC	AGACGGCATT	TCCGACGTTA
	401	TGCCGTCTGA	ACGTTCCGAT	ACTtccgcCG	AAACCCCTTG	GGACACGAGG
25	451	CAAACCGCGA	GTACAGCCGA	ACCTGCGGAA	accGACAAAG	ACCCGCGATT
	501	GCGGGAATAC	CTGactgctt	ctgccgcgcgc	acctgtctgta	Cagagggcgc
	551	tcgaagtctg	ctaTATCGAT	ACTGCTGTTG	AAACgcctgtT	tccgcacacc
	601	acttccctgc	gcaAACACGGC	AATAAACCCG	AAACCGCGATT	TttgtccgaA
	651	ACACCGCGCc	aAACCGAAat	tgcacgctcG	TAAATCATAA	

30

1	MFDFGLGELI	FVGIIALIVL	GPERLPEAAR	TAGRLIGRLQ	RFVGSVKQEL
51	DTQIELEELR	KVKQAFEEEE	AQVRDSLKET	DTDMQNSLHD	ISDGLKPWEK
101	LPEQRTPADF	GVDENGNPLP	DTANTVSDGI	SDVMPESERS	TSAETLGDDR
151	QTGSTAEPAE	TKDRAWREY	LTASAAAPVV	QRAVEVSYID	TAVETVPVHT
201	TSLRKQAVIN	KRDFCPKHRA	KPKLRVRKS*		

35	orf2.pep	MXDFLGLGELVFVGIIALIVLGPERXPEAARXAGRLIGRLQRFGVGSVKQEFDTQIELEELR	60
	orf2ng	MFDLGLGELIFVGIIALIVLGPRLPEAARTAGRLIGRLQRFGVGSVKQELDQIELEELR	60
40	orf2.pep	KAKQEFEAAAAQVRDSLKETGTDMEGNLHDISDGLKPWEKLPEQRTPADFGVDENGNPXS	120
	orf2ng	KVKQAFAEAAAAQVRDSLKETDTDMQNSLHDISDGLKPWEKLPEQRTPADFGVDEKGNSLP	120
	orf2.pep	RCGKHPIRRHFRRYAV	136
45	orf2ng	RYGKHRIRRHFRYAV	136

		10	20	30	40	50	60
50	orf2-1.pep	MFDFGLGELVFVGIIALIVLGP	ERLPEAARTAGRLIGRLQR	FVGSVKQEF	FD	TQIELEELR	
		: :					
	orf2ng-1	MFDFGLGELIFVGIIALIVLGP	ERLPEAARTAGRLIGRLQR	FVGSVKQEL	DT	TQIELEELR	
		10	20	30	40	50	60
55		70	80	90	100	110	120
	orf2-1.pep	KAKQEF	AAAAQVRDSLKETG	TDMEGN	LHDISDGLKP	WEKLPEQRT	PADFGVDENG
		: :					
	orf2ng-1	KVKQAF	AAAAQVRDSLKETD	TDMONSLH	DISDGLKP	WEKLPEQRT	PADFGVDENG
		: :					

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		70	80	90	100	110	120
		130	140	150	160	170	180
5	orf2-1.pep	DAANTLS	DGISDV	MPSE	RSYAS	AE	TLGDSGQTGSTAEPAETDQDRAWREYLTASAAAPVV
	orf2ng-1	DTANTV	SDGISD	VMPSER	SDTSA	ETLGDD	RQTGSTAEPAETDKDRAWREYLTASAAAPVV
		130	140	150	160	170	180
10	orf2-1.pep	Q-TVEV	SYIDTA	VETFP	VPH	HTSLR	KQAI
	orf2ng-1	QRAVEV	SYIDTA	VETFP	VPH	HTSLR	KQAINRKRDFCPK
		190	200	210	220	229	

Computer analysis of these amino acid sequences indicates a transmembrane region (underlined),
 15 and also revealed homology (59% identity) between the gonococcal sequence and the TatB protein
 of *E.coli*:

gnl|PID|e1292181 (AJ005830) TatB protein [Escherichia coli] Length = 171
 Score = 56.6 bits (134), Expect = 1e-07
 Identities = 30/88 (34%), Positives = 52/88 (59%), Gaps = 1/88 (1%)
 20
 Query: 1 MFD FGLGELIFVGIIALIVLGPRLPEAARTAGRLIGRLQRFVGSVKQELDTQIELEELR 60
 MFD G EL+ V II L+VLGP+RLP A +T I L+ +V+ EL +++L+E +
 Sbjct: 1 MFDIGFSELLLVFIIGLVLPQRLPVAVKTVAGWIRALRSLATTQNELTQELKLQEFQ 60
 25
 Query: 61 -KVKQAFEAAAAQVRDSLKETDMDQNS 87
 +K+ +A+ + LK + +++ +
 Sbjct: 61 DSLKKVEKASLTNLTPELKASMDLRQA 88

Based on this analysis, it was predicted that ORF2, ORF2a and ORF2ng are likely to be membrane
 proteins and so the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be
 30 useful antigens for vaccines or diagnostics, or for raising antibodies.

ORF2-1 (16kDa) was cloned in pET and pGex vectors and expressed in *E.coli*, as described above.
 The products of protein expression and purification were analyzed by SDS-PAGE. Figure 3A
 shows the results of affinity purification of the GST-fusion protein, and Figure 3B shows the results
 of expression of the His-fusion in *E.coli*. Purified GST-fusion protein was used to immunise mice,
 35 whose sera were used for Western blots (Figure 3C), ELISA (positive result), and FACS analysis
 (Figure 3D). These experiments confirm that ORF37-1 is a surface-exposed protein, and that it is
 a useful immunogen.

Example 10

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 77>:

40 1 ATGCAAGCAC GGCTGCTGAT ACCTATTCTT TTTTCAGTTT TTATTTTATC
 51 CGC.TGCGGG AACTGACAG GTATTCCATC GCATGGCGgA GkTAAACgCT
 101 TTgCGGTCGA ACAAGAACTT GTGGCCGCTT CTGCCAGAGC TGCCGTTAAA
 151 GACATGGATT TACAGGCATT ACACGGACGA AAAGTTGCAT TGTACATTGC
 201 CACTATGGGC GACCAAGGTT CAGGcAGTTT GACAGGGGGG TCGCTACTCC
 45 251 ATTGATGCAC kGrTwCstGG CGAATACATA AACAGCCCTG CCGTCCGTAC
 301 CGATTACACC TATCCACGTT ACGAAACCAC CGCTGAAACA ACATCAGGCG
 351 GTTTGACAGG TTTAACCCT TCTTTATCTA CACTTAATGC CCCTGCACTC
 401 TCTCGCACCC AATCAGACGG TAGCGGAAGT AAAAGCAGTC TGGGCTTAAA
 451 TATTGGCGGG ATGGGGGATT ATCGAAATGA AACCTTGACG ACTAACCCGC

501 GCGACACTGC CTTTCTTTCC CACTTGGTAC AGACCGTATT TTTCCTGCGC
 551 GGCATAGACG TTGTTTCTCC TGCCAATGCC GATACAGATG TGTTTATTAA
 601 CATCGACGTA TTCGGAACGA TACGCAACAG AACCAGAAATG..

This corresponds to the amino acid sequence <SEQ ID 78; ORF15>:

5 1 MQARLLIPIL FSVFILLSACG TLTGIPSHGG XKRFAVEQEL VAASARAANK
 51 DMDLQALHGR KVALYIATMG DQSGSGLTGG RYSDAXXG EYINSPAVRT
 101 DYTYPREYET AETTSGLTGT LTSLSTLNA PALSRQSDG SGSKSSLGLN
 151 IGGMDYRNE TLTNPRDTA FLHLVQTVF FLRGIDVSP ANADTDVFIN
 201 IDVFGTIRNR TEM..

10 Further work revealed the complete nucleotide sequence <SEQ ID 79>:

1 ATGCAAGCAC GGCTGCTGAT ACCTATTCTT TTTTCAGTTT TTATTTTATC
 51 CGCCTGCGGG AACTGACAG GTATTCCATC GCATGGCGGA GGTAAACGCT
 101 TTGCGGTCTGA ACAAGAACTT GTGGCCGCTT CTGCCAGAGC TGCCGTTAAA
 151 GACATGGATT TACAGGCATT ACACGGACGA AAAGTTGCAT TGTACATTGC
 15 201 CACTATGGGC GACCAAGGTT CAGGCAGTTT GACAGGGGGT CGCTACTCCA
 251 TTGATGCACT GATTCTGTGC GAATACATAA ACAGCCCTGC CGTCCGTACC
 301 GATTACACCT ATCCACGTTA CGAAACCACC GCTGAAACAA CATCAGGCGG
 351 TTTGACAGGT TTAACCACTT CTTTATCTAC ACTTAATGCC CCTGCACTCT
 401 CTGCAACCCA ATCAGACGGT AGCGGAAGTA AAAGCAGTCT GGGCTTAAAT
 20 451 ATTGGCGGGA TGGGGGATTA TCGAAATGAA ACCTTGACGA CTAACCCGCG
 501 CGACACTGCC TTTCTTTCCC ACTTGGTACA GACCGTATTT TTCCTGCGCG
 551 GCATAGACGT TGTTTCTCCT GCCAATGCCG ATACAGATGT GTTTATTAAC
 601 ATCGACGTAT TCGGAACGAT ACGCAACAGA ACCGAAATGC ACCTATACAA
 651 TGCCGAAACA CTGAAAGCCC AAACAAAAC GGAATATTTT GCAGTAGACA
 25 701 GAACCAATAA AAAATTGCTC ATCAAACCAA AAACCAATGC GTTTGAAGCT
 751 GCCTATAAAG AAAATTACGC ATTGTGGATG GGGCCGTATA AAGTAAGCAA
 801 AGGAATTAAA CCGACGGAAG GATTAATGGT CGATTTCTCC GATATCCGAC
 851 CATACGGCAA TCATACGGGT AACTCCGCCC CATCCGTAGA GGCTGATAAC
 901 AGTCATGAGG GGTATGGATA CAGCGATGAA GTAGTGGAC AACATAGACA
 30 951 AGGACAACCT TGA

This corresponds to the amino acid sequence <SEQ ID 80; ORF15-1>:

1 MQARLLIPIL FSVFILLSACG TLTGIPSHGG GKRFAVEQEL VAASARAANK
 51 DMDLQALHGR KVALYIATMG DQSGSGLTGG RYSDALIRG EYINSPAVRT
 101 DYTYPREYET AETTSGLTGT LTSLSTLNA PALSRQSDG SGSKSSLGLN
 35 151 IGGMDYRNE TLTNPRDTA FLHLVQTVF FLRGIDVSP ANADTDVFIN
 201 IDVFGTIRNR TEMHLYNAET LKAQTKLEYF AVDRTNKKLL IKPKTNAFEA
 251 AYKENYALWM GPYKVSIGIK PTEGLMVDFS DIRPYGNHTG NSAPSVEADN
 301 SHEGGYSDS VVRQHRQGP *

Further work identified the corresponding gene in strain A of *N.meningitidis* <SEQ ID 81>:

1 ATGCAAGCAC GGCTGCTGAT ACCTATTCTT TTTTCAGTTT TTATTTTATC
 51 CGCCTGCGGG AACTGACAG GTATTCCATC GCATGGCGGA GGTAAACGCT
 101 TTGCGGTCTGA ACAAGAACTT GTGGCCGCTT CTGCCAGAGC TGCCGTTAAA
 151 GACATGGATT TACAGGCATT ACACGGACGA AAAGTTGCAT TGTACATTGC
 45 201 AACTATGGGC GACCAAGGTT CAGGCAGTTT GACAGGGGGT CGCTACTCCA
 251 TTGATGCACT GATTCTGTGC GAATACATAA ACAGCCCTGC CGTCCGTACC
 301 GATTACACCT ATCCACGTTA CGAAACCACC GCTGAAACAA CATCAGGCGG
 351 TTTGACAGGT TTAACCACTT CTTTATCTAC ACTTAATGCC CCTGCACTCT
 401 CGCGCACCCA ATCAGACGGT AGCGGAAGTA AAAGCAGTCT GGGCTTAAAT
 451 ATTGGCGGGA TGGGGGATTA TCGAAATGAA ACCTTGACGA CTAACCCGCG
 50 501 CGACACTGCC TTTCTTTCCC ACTTGGTACA GACCGTATTT TTCCTGCGCG
 551 GCATAGACGT TGTTTCTCCT GCCAATGCCG ATACGGATGT GTTTATTAAC
 601 ATCGACGTAT TCGGAACGAT ACGCAACAGA ACCGAAATGC ACCTATACAA
 651 TGCCGAAACA CTGAAAGCCC AAACAAAAC GGAATATTTT GCAGTAGACA
 701 GAACCAATAA AAAATTGCTC ATCAAACCAA AAACCAATGC GTTTGAAGCT
 55 751 GCCTATAAAG AAAATTACGC ATTGTGGATG GGACCGTATA AAGTAAGCAA
 801 AGGAATTAAA CCGACAGAAG GATTAATGGT CGATTTCTCC GATATCCAAC
 851 CATACGGCAA TCATATGGGT AACTCTGCCC CATCCGTAGA GGCTGATAAC
 901 AGTCATGAGG GGTATGGATA CAGCGATGAA GCAGTGGAC GACATAGACA
 951 AGGGCAACCT TGA

60 This encodes a protein having amino acid sequence <SEQ ID 82; ORF15a>:

1 MQARLLIPIL FSVFILLSACG TLTGIPSHGG GKRFAVEQEL VAASARAANK

51 DMDLQALHGR KVALYIATMG DQSGSGLTGG RYSIDALIRG EYINSPAVRT
 101 DYTYPYR YETT AETTSGGLTG LTTSLSLTLNA PALSRTQSDG SGSKSSLGLN
 151 IGGMGDYRNE TLTTNPRDTA FLSHLVQTVF FLRGIDVVSF ANADTDVFIN
 201 IDVFGTIRNR TEMHLYNAET LKAQTKLEYF AVDRTNKKLL IKPKTNAFEA
 251 AYKENYALWM GPYKVSQGIK PTEGLMVDFS DIQPYGNHMG NSAPSVEADN
 301 SHEGYGYSDE AVRRHRQGP *

The originally-identified partial strain B sequence (ORF15) shows 98.1% identity over a 213aa overlap with ORF15a:

10	orf15.pep	10 20 30 40 50 60	MQARLLIPILFSVFILSACGTLTGIPSHGGKKRFAVEQELVAASARA AVKMDLQALHGR
	orf15a	10 20 30 40 50 60	MQARLLIPILFSVFILSACGTLTGIPSHGGKKRFAVEQELVAASARA AVKMDLQALHGR
15	orf15.pep	70 80 90 100 110 120	KVALYIATMGDQSGSGLTGGRYSIDAXXGEYINSPAVRTDYTYPRYETTAETTSGGLTG
	orf15a	70 80 90 100 110 120	KVALYIATMGDQSGSGLTGGRYSIDALIRGEYINSPAVRTDYTYPRYETTAETTSGGLTG
20	orf15.pep	130 140 150 160 170 180	LTTSLSLTLNAPALSRTQSDGSGSKSSLGLNIGMGDYRNETLTNPRDTAFLSHLVQTVF
	orf15a	130 140 150 160 170 180	LTTSLSLTLNAPALSRTQSDGSGSKSSLGLNIGMGDYRNETLTNPRDTAFLSHLVQTVF
25	orf15.pep	190 200 210	FLRGIDVVS PANADTDVFINIDVFGTIRNRTEM
	orf15a	190 200 210 220 230 240	FLRGIDVVS PANADTDVFINIDVFGTIRNRTEMHLYNAETLKAQTKLEYFAVDRTNKKLL

The complete strain B sequence (ORF15-1) and ORF15a show 98.8% identity in 320 aa overlap:

35	orf15a.pep	10 20 30 40 50 60	MQARLLIPILFSVFILSACGTLTGIPSHGGKKRFAVEQELVAASARA AVKMDLQALHGR
	orf15-1	10 20 30 40 50 60	MQARLLIPILFSVFILSACGTLTGIPSHGGKKRFAVEQELVAASARA AVKMDLQALHGR
40	orf15a.pep	70 80 90 100 110 120	KVALYIATMGDQSGSGLTGGRYSIDALIRGEYINSPAVRTDYTYPRYETTAETTSGGLTG
	orf15-1	70 80 90 100 110 120	KVALYIATMGDQSGSGLTGGRYSIDALIRGEYINSPAVRTDYTYPRYETTAETTSGGLTG
45	orf15a.pep	130 140 150 160 170 180	LTTSLSLTLNAPALSRTQSDGSGSKSSLGLNIGMGDYRNETLTNPRDTAFLSHLVQTVF
	orf15-1	130 140 150 160 170 180	LTTSLSLTLNAPALSRTQSDGSGSKSSLGLNIGMGDYRNETLTNPRDTAFLSHLVQTVF
50	orf15a.pep	190 200 210 220 230 240	FLRGIDVVS PANADTDVFINIDVFGTIRNRTEMHLYNAETLKAQTKLEYFAVDRTNKKLL
	orf15-1	190 200 210 220 230 240	FLRGIDVVS PANADTDVFINIDVFGTIRNRTEMHLYNAETLKAQTKLEYFAVDRTNKKLL
55	orf15a.pep	250 260 270 280 290 300	IKPKTNAFEAAAYKENYALWMGPYKVSQGIKPTTEGLMVDFS DIQPYGNHMGNSAPSVEADN
	orf15-1	250 260 270 280 290 300	IKPKTNAFEAAAYKENYALWMGPYKVSQGIKPTTEGLMVDFS DIRPYGNHMGNSAPSVEADN
60	orf15a.pep	310 320	SHEGYGYSDEAVRRHRQGPX
	orf15-1	310 320	SHEGYGYSDEVVRQHRQGPX

310

320

Further work identified the corresponding gene in *N.gonorrhoeae* <SEQ ID 83>:

```

      1 ATGCGGGGAC GGCTGCTGAT ACCTATTCTT TTTTCAGTTT TTATTTTATC
      51 CGCCTGCGGG ACACTGACAG GTATTCCATC GCATGGCGGA GGCAAACGCT
5   101 TCGCGGTGCGA ACAAGAACTT GTGGCCGCTT CTGCCAGAGC TGCCGTAAAA
      151 GACATGGATT TACAGGCATT ACACGGACGA AAAGTTGCAT TGTACATTGC
      201 AACTATGGGC GACCAAGGTT CAGGCAGTTT GACAGGGGGT CGCTACTCCA
      251 TTGATGCACT GATTGCGGGC GAATACATAA ACAGCCCTGC CGTCCGCACC
      301 GATTACACCT ATCCGCGTTA CGAAACCACC GCTGAAACAA CATCAGGCGG
10  351 TTTGACGGGT TTAACCACTT CTTTATCTAC ACTTAATGCC CCTGCACTCT
      401 CGCGCACCCA ATCAGACGGT AGCGGAAGTA GGAGCAGTCT GGGCTTAAAT
      451 ATTGGCGGGA TGGGGGATTA TCGAAATGAA ACCTTGACGA CCAACCCGCG
      501 CGACACTGCC TTCTTTTCCC ACTTGGTGCA GACCGTATTT TTCCTGCGCG
      551 GCATAGACGT TGTTTCTCCT GCCAATGCCG ATACAGATGT GTTTATTAAC
15  601 ATCGACGTAT TCGGAACGAT ACGCAACAGA ACCGAAATGC ACCTATACAA
      651 TGCCGAAACA CTGAAAGCCC AAACAAACT GGAATATTTC GCAGTAGACA
      701 GAACCAATAA AAAATTGCTC ATCAAACCCA AAACCAATGC GTTTGAAGCT
      751 GCCTATAAAG AAAATTACGC ATTGTGGATG GGGCCGTATA AAGTAAGCAA
      801 AGGAATCAAA CCGACGGAAG GATTGATGGT CGATTTCCTC GATATCCAAC
20  851 CATACGGCAA TCATACGGGT AACTCCGCCC CATCCGTAGA GGCTGATAAC
      901 AGTCATGAGG GGTATGGATA CAGCGATGAA GCAGTGCAC AACATAGACA
      951 AGGGCAACCT TGA

```

This encodes a protein having amino acid sequence <SEQ ID 84; ORF15ng>:

```

      1 MRARLLIPIL FSVFILSACG TLTGIPSHGG GKRFQVEQEL VAASARAAVK
25  51 DMDLQALHGR KVALYIATMG DQSGSLTGG RYSIDALIRG EYINSPAVRT
      101 DYTYPYRSET AETTSGLTGT LTSLSTLNA PALSRTQSDG SGRSSLGLN
      151 IGGMGDYRNE TLTNPRDTA FLSHLVQTVF FLRGIDVSP ANADTDVFIN
      201 IDVFGTIRNR TEMHLYNAET LKAQTKLEYF AVDRTNKKLL IKPKTNAFEA
      251 AYKENYALWM GPYKVSCKIK PTEGLMVDFS DIQPYGNHTG NSAPSVEADN
30  301 SHEGYGYSDE AVRQHRQGP *

```

The originally-identified partial strain B sequence (ORF15) shows 97.2% identity over a 213aa overlap with ORF15ng:

```

      orf15.pep      MQARLLIPILFSVFI LSACGTLTGIPSHGGGKRFQVEQELVAASARA AVKMDMDLQALHGR      60
35  orf15ng          MRARLLIPILFSVFI LSACGTLTGIPSHGGGKRFQVEQELVAASARA AVKMDMDLQALHGR      60
      orf15.pep      KVALYIATMGDQSGSLTGGRYSIDAXXGEYINSPAVRTDYTYPRYETTAETTSGLTGT      120
40  orf15ng          KVALYIATMGDQSGSLTGGRYSIDALIRGEYINSPAVRTDYTYPRYETTAETTSGLTGT      120
      orf15.pep      LTSLSTLNA PALSRTQSDGSGSKSSLGLNIGMGDYRNETLTNPRDTAFLSHLVQTVF      180
      orf15ng          LTSLSTLNA PALSRTQSDGSGSRSSLGLNIGMGDYRNETLTNPRDTAFLSHLVQTVF      180
45  orf15.pep      FLRGIDVVS PANADTDVFINIDVFGTIRNRTEM      213
      orf15ng          FLRGIDVVS PANADTDVFINIDVFGTIRNRTEMHLYNAETLKAQTKLEYFAVDRTNKKLL      240

```

The complete strain B sequence (ORF15-1) and ORF15ng show 98.8% identity in 320 aa overlap:

```

50  orf15-1.pep      10      20      30      40      50      60
      orf15ng          10      20      30      40      50      60
      orf15-1.pep      MQARLLIPILFSVFI LSACGTLTGIPSHGGGKRFQVEQELVAASARA AVKMDMDLQALHGR
      orf15ng          MRARLLIPILFSVFI LSACGTLTGIPSHGGGKRFQVEQELVAASARA AVKMDMDLQALHGR
55  orf15-1.pep      70      80      90      100     110     120
      orf15ng          70      80      90      100     110     120
      orf15-1.pep      KVALYIATMGDQSGSLTGGRYSIDALIRGEYINSPAVRTDYTYPRYETTAETTSGLTGT
      orf15ng          KVALYIATMGDQSGSLTGGRYSIDALIRGEYINSPAVRTDYTYPRYETTAETTSGLTGT
60  orf15-1.pep      130     140     150     160     170     180
      orf15ng          130     140     150     160     170     180
      orf15-1.pep      LTSLSTLNA PALSRTQSDGSGSKSSLGLNIGMGDYRNETLTNPRDTAFLSHLVQTVF

```

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```

      |||
orf15ng      |||:|||||
              130      140      150      160      170      180
5      orf15-1.pep      190      200      210      220      230      240
      orf15ng      190      200      210      220      230      240
10     orf15-1.pep      250      260      270      280      290      300
      orf15ng      250      260      270      280      290      300
15     orf15-1.pep      310      320
      orf15ng      310      320
20     orf15-1.pep      SHEGYGYSDEVVRQHRQGQPX
      orf15ng      SHEGYGYSDEAVRQHRQGQPX
              310      320

```

Computer analysis of these amino acid sequences reveals an ILSAC motif (putative membrane lipoprotein lipid attachment site, as predicted by the MOTIFS program).

indicates a putative leader sequence, and it was predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

ORF15-1 (31.7kDa) was cloned in pET and pGex vectors and expressed in *E.coli*, as described above. The products of protein expression and purification were analyzed by SDS-PAGE. Figure 4A shows the results of affinity purification of the GST-fusion protein, and Figure 4B shows the results of expression of the His-fusion in *E.coli*. Purified GST-fusion protein was used to immunise mice, whose sera were used for Western blot (Figure 4C) and ELISA (positive result). These experiments confirm that ORFX-1 is a surface-exposed protein, and that it is a useful immunogen.

Example 11

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 85>:

```

35      1      ..GG.CAGCACA AAAACAGGC GGTGAACGG AAAAACCGTA TTTACGATGA
      51      TGCCGGGTAT GATATTCGGC GTATTCACGG GCGCATTCTC CGCAAAATAT
      101     ATCCCCGCGT TCGGGCTTCA AATTTTCTTC ATCCTGTTTT TAACCGCCGT
      151     CGCATTCAA AACTGCATA CCGACCCTCA GACGGCATCC CGCCCGCTGC
      201     CCGGACTGCC CCGACTGACT GCGGTTTCCA CACTGTTCCG CACAATGTCG
40     251     AGCTGGGTGC GCATAGGCGG CGGTTCACTT TCCGTCCCCT TCTTAATCCA
      301     CTGCGGCTTC CCGGCCATA AAGCCATCGG CACATCATCC GGCCTTGCCCT
      351     GGCCGATTGC ACTCTCCGGC GCAATATCGT ATCTGCTCAA CGGCCTGAAT
      401     ATTGCAGGAT TGCCCGAAGG GTCACTGGGC TCCTTTTACC TGCCCGCCGT
      451     CGCCGTCTTC AGCGCGGCAA CCATTGCCTT TGCCCGCTC GGTGTCAAAA
45     501     CCGCCACAA ACTTTCTTCT GCCAACTCA AAAAATC.TT CGGCATTATG
      551     TTGCTTTTGA TTGCCGAAA AATGCTGTAC AACCTGCTTT AA

```

This corresponds to the amino acid sequence <SEQ ID 86; ORF17>:

```

      1      ..GQHKQAVNG KTVFTMMPGM IFGVFTGAFS AKYIPAFGLQ IFFILFLTAV
      51      AFKTLHTDPQ TASRPLPLP XLTAVSTLFG TMSSWVGIGG GSLSVPFLLH

```

5	1	ATGTGGCATT	GGGACATTAT	CTTAATCCTG	CTTGCCGTAG	GCAGTGCGCC
	51	AGGTTTTATT	GCCGGCCTGT	TCGGCGTAGG	CGCGCGCACG	CTGATTGTCC
	101	CTGTCGTTTT	ATGGGTGCTT	GATTTGCAGG	GTTTGGCACA	ACATCCTTAC
	151	CGCACAACCC	TCGCGTCCGG	CACATCCTTC	GGCGTCATGG	TCTTCACCGC
10	201	CTTTTCCAGT	ATGCTGGGGC	AGCACAAAAA	ACAGGGCGGT	GACTGGAAAA
	251	CCGTATTTAC	GATGATGCCG	GGTATGATAT	TCGGCGTATT	CACGGGCGCA
	301	CTCTCCGCAA	AATATATCCC	CGCGTTCGGG	CTTCAAAATT	TCTTCATCCT
	351	GTTTTTAAAC	GCCGTGCGAT	TCAAAACACT	GCATACCGAC	CCTCAGACGG
15	401	CATCCGCCCC	GCTGCCCGGA	CTGCCCGGAC	TGACTCGGGT	TCTCCACACT
	451	TTCCGGCACA	TGTCGAGCTG	GGTCGGCATA	GGCGGCGGTT	TTACCTCCGT
	501	CCCCTTCTTA	ATCCACTGCG	GCTTCCCCCG	CCATAAAGCC	ATCGGCACAT
	551	CATCCGGCCT	TGCCTGGCCG	ATTGCACTCT	CCGGGCGCAAT	ATCGTATCTG
20	601	CTCAACGGCC	TGAATTATTG	AGGATTGCCC	GAAAGGTCAC	TGGGCTTCTT
	651	TTACCTTGCCC	GGCGTCGCCG	TCCTCAGCGC	GGACAACATT	GCCTTTGCCC
	701	CGCTCGGTGT	CAAAACCGCC	CACAAACTTT	CTTCTGCCAA	ACTCAAAAAA
	751	Tc.TTCGGCA	TTATGTTGCT	TTTGATTGCC	GGAAAAATGC	TGTACAACCT
	801	GCTTTTAA				

25

1	<u>MWHWDIILIL</u>	<u>LAVGSAAGFI</u>	<u>AGLFGVGGGT</u>	<u>LIVPVVLWVL</u>	<u>DLQGLAQHPY</u>
51	<u>LAQHLAVGTST</u>	<u>AVMVFTAFSS</u>	<u>MLGQHKQAV</u>	<u>PDKTVFTMPP</u>	<u>GMIFGVFTGA</u>
101	<u>LSAKHYPAFG</u>	<u>LQIIFLFLT</u>	<u>AVAEKTLHTD</u>	<u>PQTASRELPG</u>	<u>LPGLATVSTA</u>
151	<u>FGTMSSWVGI</u>	<u>GGGSLSVFFL</u>	<u>THCGFPAHKA</u>	<u>IGTSSGLAWP</u>	<u>IALSGAISYL</u>
201	<u>LNGLNIAGLP</u>	<u>EGSLGLFLYP</u>	<u>AVAVLSAATI</u>	<u>AFAPLGVKTA</u>	<u>HKLSSAKLKK</u>
251	<u>XFGIMLLLIA</u>	<u>GKMLYNLL*</u>			

Homology with hypothetical *H.influenzae* transmembrane protein HI0902 (accession number P44070)

	ORF17	3	HKKQAVNGKTVFTMMPGMIFGVFT-GAFSAKYIPAFGLQIF--FILFLTAVAFKTLHTDP	59
			HK + + V + P ++ VF G F + +IF +++L ++ D	
	HI0902	72	HKLGNIVWQAVRILAPVIMLSVFICGLFIGRLDREISAKIFACLVVYLATKMVLSIKKD-	130
35	ORF17	60	QTASRPLPGLPXLTAVSTLFGTMSSWVGIGGGSLSPVFLIHCGFPAHKAIGTSSGLAWPI	119
			Q ++ L L + L G SS GIGGG VPFL G +AIG+S+ +	
	HI0902	131	QVTTKSLTPLSSVIG-GILIGMASSAAGIGGGGFIVPFLTARGINIKQAIGSSAFCGMLL	189
40	ORF17	120	ALSGAISYLLNGLNIAGLPEGSLGFLYLPAAVAVLSAATIAFAPLVGXXXXXXXXXXXXXXXXX	179
			+SG S+++G +PE SLG++YLPV ++A + + LG	
	HI0902	190	GISGMFSFIVSGWGNPLMPEYSLGYIYLPVAVLGITATSFFTSLKGASATAKLPVSTLKKG	249
	ORF17	180	FGIMLLLIAGKM	191
			F + L+++A M	
45	HI0902	250	FALFLIVVAINM	261

ORF17 shows 96.9% identity over a 196aa overlap with an ORF (ORF17a) from strain A of *N. meningitidis*:

```

50                                     10      20      30
    orf17.pep                        GQHKKQAVNGKTVFTMMPGMIFGVFTGAFS
                                     |||||: |||||: |||||: |||||
    orf17a      QGLAQHPYAQHLAVGTSFAVMVFTAFSSMLGQHKKQAVDWKTVFTMMPGMVFGVFAGALS
                50      60      70      80      90      100

55                                     40      50      60      70      80      90
    orf17.pep      AKYIPAFGLQIFFILFLTAVAFKTLHTDPQTASRPLPGLPXLTAVSTLFGTMSSWVGIGG
                   ||||| ||||| ||||| ||||| ||||| |||||
    orf17a      AKYIPAFGLQIFFILFLTAVAFKTLHTDPQTASRPLPGLPGLTAVSTLFGTMSSWVGIGG

```

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		110	120	130	140	150	160
		100	110	120	130	140	150
5	orf17.pep	GSLSVPF	IHC	GFP	PAHKA	IGTSS	GLAWPIALSGAIS
	orf17a	GSLSVPF	IHC	GFP	PAHKA	IGTSS	GLAWPIALSGAIS
		170	180	190	200	210	220
10		160	170	180	190		
	orf17.pep	AVLSAAT	IAFAP	LG	VKT	AHKLSSAKL	KKSF
	orf17a	AVLSAAT	IAFAP	LG	VKT	AHKLSSAKL	KKSF
		230	240	250	260		

The complete length ORF17a nucleotide sequence <SEQ ID 89> is:

15	1	ATGTGGCATT	GGGACATTAT	CTTAATCCTG	CTTGCCGTAG	GCAGTGC	CGGC
	51	AGGTTTTTATT	GCCGGCCTGT	TCGGCGTAGG	CGGCGGCACG	CTGATTGTCC	
	101	CTGTCGTTTT	ATGGGTGCTT	GATTGTCAGG	GTTTGGCACA	ACATCCTTAC	
	151	GCGCAACACC	TCGCCGTCGG	CACATCCTTC	GCCGTCATGG	TCTTCACCGC	
	201	CTTTTCCAGT	ATGCTGGGGC	AGCACAAAAA	ACAGGCGGTC	GACTGGAAAA	
20	251	CCGTATTTAC	GATGATGCCG	GGTATGGTAT	TCGGCGTATT	CGCTGGCGCA	
	301	CTCTCCGCAA	AATATATCCC	AGCGTTCGGG	CTTCAAATTT	TCTTCATCCT	
	351	GTTTTTAACC	GCCGTCGCAT	TCAAACACT	GCATACCGAC	CCTCAGACGG	
	401	CATCCCGCCC	GCTGCCCGGA	CTGCCCGGAC	TGACTGCGGT	TTCCACACTG	
	451	TTCCGCACAA	TGTCGAGCTG	GGTCGGCATA	GGCGCGGCTT	CACCTTCCGT	
25	501	CCCCTTCTTA	ATCCACTGCG	GCTTCCCCGC	CCATAAAGCC	ATCGGCACAT	
	551	CATCCGGCCT	TGCCTGGCCG	ATTGCACTCT	CCGCGCCAAT	ATCGTATCTG	
	601	CTCAACGGCC	TGAATATTGC	AGGATTGCCC	GAAGGCTCAC	TGGGCTTCCT	
	651	TTACCTGCCC	GCCGTCGCCG	TCCTCAGCGC	GGCAACCATT	GCCTTTGCCC	
	701	CGCTCGGTGT	CAAAACCGCC	CACAACTTT	CTTCTGCCAA	ACTCAAAAAA	
30	751	TCCTTCGGCA	TTATCTTGCT	TTTGATTGCC	GGAAAAATGC	TGTACAACCT	
	801	GCTTTAA					

This encodes a protein having amino acid sequence <SEQ ID 90>:

	1	MWHWDIILIL	LAVGSAAGFI	AGLFGVGGGT	LIVPVVLWVL	DLQGLAQHPY	
	51	AQHLAVGTSF	AVMVFTAFSS	MLGQHKQAV	DWKTVFTMMP	GMVFGVFAGA	
35	101	LSAKYIPAFG	LQIFFILFLT	AVAFKTLHTD	PQTASRPLPG	LPGLTAVSTL	
	151	FGTMSSWVG	IGGSLSVFPL	IHC	GFP	PAHKA	IGTSSGLAWP
	201	LNGLNIAGLP	EGSLGFLYLP	AVAVLSAATI	AFAPLGVKTA	HKLSSAKLKK	
	251	SFGIMLLLIA	GKMLYNLL*				

ORF17a and ORF17-1 show 98.9% identity in 268 aa overlap:

40		10	20	30	40	50	60
	orf17a.pep	MWHWDIILIL	LAVGSAAGFI	AGLFGVGGGT	LIVPVVLWVL	DLQGLAQHPY	AQHLAVGTSF
	orf17-1						
45		10	20	30	40	50	60
	orf17a.pep	AVMVFTAFSS	MLGQHKQAV	DWKTVFTMMP	GMVFGVFAG	ALSAKYIPAF	GLQIFFILFLT
	orf17-1						
50		70	80	90	100	110	120
	orf17a.pep	AVMVFTAFSS	MLGQHKQAV	DWKTVFTMMP	GMIFGVFTG	ALSAKYIPAF	GLQIFFILFLT
	orf17-1						
55		130	140	150	160	170	180
	orf17a.pep	AVAFKTLHTD	PQTASRPLPG	LPGLTAVSTL	FGTMSSWVG	IGGSLSVFPL	IHC
	orf17-1						
60		130	140	150	160	170	180
	orf17a.pep	IGTSSGLAWPI	ALSGAISYLL	NGLNIAGL	PEGSLGFLYLP	PAVAVLSAATI	AFAPLGVKTA
	orf17-1						
65		190	200	210	220	230	240
	orf17a.pep	IGTSSGLAWPI	ALSGAISYLL	NGLNIAGL	PEGSLGFLYLP	PAVAVLSAATI	AFAPLGVKTA
	orf17-1						
		190	200	210	220	230	240
	orf17a.pep	HKLSSAKLKK	SFGIMLLLIA	GKMLYNLLX			
		250	260	269			

```

      |||||
orf17-1      HKLSSAKLKKXFGIMLLLIAGKMLYNLLX
              250      260

```

5 Homology with a predicted ORF from *N.gonorrhoeae*

ORF17 shows 93.9% identity over a 196aa overlap with a predicted ORF (ORF17.ng) from *N. gonorrhoeae*:

```

      orf17.pep                                GQHKKQAVNGKTVFTMPGMIFGVFTGAFS      30
10  orf17ng      QGLAQHPYAQHLAVGTSFAVMVFTAFSSMLGQHKKQAVDWKTIFAMMPGMIFGVFAGALS      102
      orf17.pep      AKYIPAFGLQIFFILFLTAVAFKTLHTDPQTASRPLPGLPXLTAVSTLFGTMSSWVGIGG      90
      orf17ng      AKYIPAFGLQIFFILFLTAVAFKTLHTGRQTASRPLPGLPGLTAVSTLFGAMSSWVGIGG      162
15  orf17.pep      GSLSVPFLIHCGFPAHKAIGTSSGLAWPIALSGAISYLLNGLNIAGLPEGSLGFLYLPVAV      150
      orf17ng      GSLSVPFLIHCGFPAHKAIGTSSGLAWPIALSGAISYLVNGLNIAGLPEGSLGFLYLPVAV      202
20  orf17.pep      AVLSAATIAFAPLGVKTAHKLSSAKLKSFGIMLLLIAGKMLYNLL      196
      orf17ng      AVLSAATIAFAPLGVKTAHKLSSAKLKESFGIMLLLIAGKMLYNLL      268

```

An ORF17ng nucleotide sequence <SEQ ID 91> is predicted to encode a protein having amino acid sequence <SEQ ID 92>:

```

25      1  MWHWDIILIL LAVGSAAGFI AGLFGVGGGT LIVPVVLWVL DLQGLAQHPY
      51  AQHLAVGTSF AVMVFTAFSS MLGQHKKQAV DWKTIFAMMP GMIFGVFAGA
      101  LSAKYIPAFG LQIFFILFLT AVAFKTLHTG RQTASRPLPG LPGLTAVSTL
      151  FGAMSSWVGI GGSLSVPFL IHCGFPAHKA IGTSSGLAWP IALSGAISYL
30  201  VNGLNIAGLP EGSLGFLYLP AVAVLSAATI AFAPLGVKTA HKLSSAKLKE
      251  SFGIMLLLIA GKMLYNLL*

```

Further work revealed the complete gonococcal DNA sequence <SEQ ID 93>:

```

      1  ATGTGGCATT GGGACATTAT CTTAATCCTG CTTGCcgtag gcAGTGCGGC
      51  AGGTTTTATT GCCGGCCTGT Tcgggtgtagg cggcgtACAG CTGATTGTCC
35  101  CTGTCGTTTT ATGGGTGCTT GATTGTCAGG GTTTGGCACA ACATCCTTAC
      151  GCGCAACACC TCGCCGTCGG CACaTccttc gcCGTCATGG TCTTCACCGC
      201  CTTTCCAGT ATGTGGGGC AGCACAAAAA ACAGGCGGTC GACTGGAAAA
      251  CCATATTTGC GATGATGCCG GGTATGATAT TCGGCGTATT CGCTGGCGCA
      301  CTCTCCGCAA AATATATCCC CGCGTTCGGG CTTCAAATTT TCTTCATCCT
      351  GTTTTTAACC GCCGTCGCAT TCAAAACACT GCATACCGGT CGTCAGACGG
40  401  CATCCCGCCC GCTGCCCGGG CTGCCCGGAC TGACTGCGGT TTCCACACTG
      451  TTCGGCGCAA TGTCGAGCTG GGTGGGCATA GCGGCGGTT CACTTTCCGT
      501  CCCCTTCTTA ATCCACTGCG GCTTCCCCGC CCATAAAGCC ATCGGCACAT
      551  CATCCGGCCT TGCCTGGCCG ATTGCACTCT CCGGCGCAAT ATCGTATCTG
      601  GTCAACGGTC TGAATATTGC AGGATTGCCG GAAGGGTCGC TGGGCTTCCT
45  651  TTACCTGCCC GCCGTCGCCG TCCTCAGCGC GGCAACCATT GCCTTTGCCC
      701  CGCTCGGTGT CAAAACCGCC CACAAACTTT CTTCTGCCAA ACTCAAAGAA
      751  TCCTTCGGCA TTATGTTGCT TTTGATTGCC GAAAAAATGC TGTACAACCT
      801  GCTTTAA

```

This corresponds to the amino acid sequence <SEQ ID 94; ORF17ng-1>:

```

50      1  MWHWDIILIL LAVGSAAGFI AGLFGVGGGT LIVPVVLWVL DLQGLAQHPY
      51  AQHLAVGTSF AVMVFTAFSS MLGQHKKQAV DWKTIFAMMP GMIFGVFAGA
      101  LSAKYIPAFG LQIFFILFLT AVAFKTLHTG RQTASRPLPG LPGLTAVSTL
      151  FGAMSSWVGI GGSLSVPFL IHCGFPAHKA IGTSSGLAWP IALSGAISYL
55  201  VNGLNIAGLP EGSLGFLYLP AVAVLSAATI AFAPLGVKTA HKLSSAKLKE
      251  SFGIMLLLIA GKMLYNLL*

```

ORF17ng-1 and ORF17-1 show 96.6% identity in 268 aa overlap:

```

              10      20      30      40      50      60
orf17-1.pep  MWHWDIILILAVGSAAGFIAGLFGVGGTLLIVPVVLWVLDLQGLAQHPYAQHLAVGTSF

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	orfl7ng-1	 MWHWDIILILLAVGSAAGFTIAGLFGVGGTLLIVPVVLWVLDLQGLAQHPYAQHLAVGTSF
		10 20 30 40 50 60
5	orfl7-1.pep	70 80 90 100 110 120 AVMVFTAFSSMLGQHKKQAVDWKTIVFTMMPGMIFGVFTGALSAKYIPAFGLQIFFILFLT
	orfl7ng-1	 AVMVFTAFSSMLGQHKKQAVDWKTIFAMMPGMIFGVFAGALSAKYIPAFGLQIFFILFLT
10		70 80 90 100 110 120
	orfl7-1.pep	130 140 150 160 170 180 AVAFKTLHTDPQTASRPLPGLPGLTAVSTLFGTMSSWVGIGGGSLSVPFLIHC GFPAHKA
	orfl7ng-1	 AVAFKTLHTGRQTASRPLPGLPGLTAVSTLFGAMSSWVGIGGGSLSVPFLIHC GFPAHKA
15		130 140 150 160 170 180
	orfl7-1.pep	190 200 210 220 230 240 IGTSSGLAWPIALSGAISYLLNGLNIAGLPEGSLGFLYLPAAVLSAATIAFAPLGVKTA
20	orfl7ng-1	 IGTSSGLAWPIALSGAISYLVNGLNIAGLPEGSLGFLYLPAAVLSAATIAFAPLGVKTA
		190 200 210 220 230 240
	orfl7-1.pep	250 260 269 HKLSSAKLKKXFGIMLLLIAGKMLYNLLX
25	orfl7ng-1	 HKLSSAKLKESFGIMLLLIAGKMLYNLLX
		250 260

In addition, ORF17ng-1 shows significant homology with a hypothetical *H. influenzae* protein:

sp|P44070|Y902_HAEIN HYPOTHETICAL PROTEIN HI0902 pir||G64015 hypothetical protein
HI0902 - Haemophilus influenzae (strain Rd KW20) gi|1573922 (U32772) H. influenzae
predicted coding region HI0902 [Haemophilus influenzae]Length = 264
Score = 74 (34.9 bits), Expect = 1.6e-23, Sum P(2) = 1.6e-23
Identities = 15/43 (34%), Positives = 23/43 (53%)

Query: 55 AVGTSFAVMVFTAFSSMLGQHKKQAVDWKTIFAMMPGMIFGVF 97
A+GTSFA +V T S HK + W+ + + P ++ VF
Sbjct: 52 ALGTSFATIVITGIGSAQRHHKLGNIWQAVRIIAPVIMLSVF 94

Score = 195 (91.9 bits), Expect = 1.6e-23, Sum P(2) = 1.6e-23
Identities = 44/114 (38%), Positives = 65/114 (57%)

Query: 150 LFGAMSSWVGIGGGSLSVPFLIHC GFPAHKAIGTSSGLAWPIALSGAISYLVNGLNIAGL 209
L G SS GIGGG VPFL G +AIG+S+ + +SG S++V+G +
Sbjct: 148 LIGMASSAAGIGGGGFIVPFLTARGINIKQAIGSSAFCGMLLGISGMFSFIVSGWGNPLM 207

Query: 210 PEGSLGFLYLPAAVLSAATIAFAPLGVKTAHKLSSAKLKESFGIMLLLIAGKM 263
PE SLG++YLPAAV ++A + + LG KL + LK+ F + L+++A M
Sbjct: 208 PEYSLGYIYLPVGLGITATSFFTSLKGASATAKLPVSTLKKGFALFLIVVAINM 261

This analysis, including the homology with the hypothetical *H. influenzae* transmembrane protein, suggests that the proteins from *N. meningitidis* and *N. gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 12

The following partial DNA sequence was identified in *N. meningitidis* <SEQ ID 95>:

1 ..GGAAACGGAT GGCAGGCAGA CCCC GAACAT CCGCTGCTCG GGCTTTTTCG
51 CGTCAGTAAT GTATCGATGA CGCTTGCTTT TGTCGGAATA TGTGCGTTGG
101 TGCATTATG CTTTTCGGGA ACGGTTCAAG TGTTTGTGTT TGCGGCACTG
151 CTCAAACCTT ATGCGCTGAA GCCGGTTTAT TGGTTCGTGT TGCAGTTTGT
201 GCTGATGGCG GTTGCCTATG TCCACCGCTG CGGTATAGAC CGGCAGCCGC
251 CGTCAACGTT CGGCGGCTCG CAGCTGCGAC TCGGCGGGTT GACGGCAGCG

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301 TTGATGCAGG TCTCGGTACT GGTGCTGCTG CTTTCAGAAA TTGGAAGATA
351 A

This corresponds to the amino acid sequence <SEQ ID 96; ORF18>:

5 1 ..GNGWQADPEH PLLGLFAVSN VSMTLAFVGI CALVHYCFSG TVQVFVFAAL
51 LKLYALKPVY WFLVQFVLMV VAYVHRCGID RQPPSTFGGS QLRLGGLTAA
101 LMQVSVLVLL LSEIGR*

Further work revealed the complete nucleotide sequence <SEQ ID 97>:

1 ATGATTTTGC TGCATTTGGA TTTTGTCT GCCTTACTGT ATGCGGCGGT
51 TTTTCTGTTT CTGATATTCC GCGCAGGAAT GTTGCAATGG TTTTGGGCGA
101 GTATTATGCT GTGGCTGGGC ATATCGGTTT TGGGGGCAAA GCTGATGCCC
151 GGCATATGGG GAATGACCCG CGCCGCGCCC TTGTTCATCC CCCATTTTAA
201 CCTGACTTTG GGCAGCATAT TTTTTCAT CGGGCATTGG AACCGGAAAA
251 CAGATGGAAA CGGATGGCAG GCAGACCCG AACATCCGCT GCTCGGGCTT
301 TTTGCCGTCA GTAATGTATC GATGACGCTT GCTTTGTGTC GAATATGTGC
15 351 GTTGGTGCA TATTGCTTTT CGGAACGGT TCAAGTGTTT GTGTTGCGG
401 CACTGCTCAA ACTTTATGCG CTGAAGCCGG TTTATTGGTT CGTGTGCGAG
451 TTTGTGCTGA TGGCGGTTGC CTATGTCCAC CGCTGCGGTA TAGACCGGCA
501 GCCGCCGTCA ACGTTCGGCG GCTCGCAGCT GCGACTCGGC GGGTTGACGG
551 CAGCGTTGAT GCAGGTCTCG GTACTGGTGC TGCTGCTTTC AGAAATTGGA
20 601 AGATAA

This corresponds to the amino acid sequence <SEQ ID 98; ORF18-1>:

1 MILHLDFLS ALLYAAVFLF LIFRAGMLQW FWASIMLWLG ISVLGAKLMP
51 GIWGMTRAAP LFIPHFYLTLSIFFFFIGHW NRKTDGNGWQ ADPEHPLLGL
101 FAVSNVSM TLAFVICALVH YCFSGTVQVF VFAALLKLYA LKPVYWFVLQ
25 151 FVLMAVAVH RCGIDRQPPS TFSGSQLRLG GLTAALMQVS VLVLLLSEIG
201 R*

Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF18 shows 98.3% identity over a 116aa overlap with an ORF (ORF18a) from strain A of *N.*

30 *meningitidis*:

				10	20	30
orf18.pep				GNGWQADPEHPLLGLFAVSNVSMTLAFVGI		
35 orf18a	TRAAPLFIPHFYLTLSIFFFFIGHWNRKTDGNGWQADPEHPLLGLFAVSNVSMTLAFVGI	60	70	80	90	100 110
		40	50	60	70	80 90
orf18.pep	CALVHYCFSGTVQVFVFAALLKLYALKPVYWFVLQFVLMAVAVHRCGIDRQPPSTFGGS					
40 orf18a	CALVHYCFSTVQVFVFAALLKLYALKPVYWFVLQFVLMAVAVHRCGIDRQPPSTFGGS	120	130	140	150	160 170
		100	110			
45 orf18.pep	QLRLGGLTAALMQVSVLVLLLSEIGRX					
orf18a	QLRLGGLTAALMQXSVLVLLLSEIGRX	180	190	200		

The complete length ORF18a nucleotide sequence <SEQ ID 99> is:

50 1 ATGATTTTGC TGCATTTGGA TTTTGTCT GCCTTACTGT ATGCGGCGGT
51 TTTTCTGTTT CTGATATTCC GCGCAGGAAT GTTGCAATGG TTTTGGGCGA
101 GTATTATGCT GTGGCTGGGC ATATCGGTTT TGGGGGCAAA GCTGATGCCC
151 GGCATATGGG GAATGACCCG CGCCGCGCCC TTGTTCATCC CCCATTTTAA
201 CCTGACTTTG GGCAGCATAT TTTTTCAT CGGGCATTGG AACCGGAAAA
251 CGGATGGAAA CGGATGGCAG GCAGACCCG AACATCCTCT GCTCGGGCTG
55 301 TTTGCCGTCA GTAATGTATC GATGACGCTT GCTTTGTGTC GAATATGTGC
351 GTTGGTGCA TATTGCTTTT CGNGAACGGT TCAAGTGTTT GTGTTGCGG
401 CACTGCTCAA ACTTTATGCG CTGAAGCCGG TTTATTGGTT CGTGTGCGAG

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451 TTTGTGCTGA TGGCGGTTGC CTATGTCCAC CGCTGCGGTA TAGACCGGCA
 501 GCCGCCGTCA ACGTTCGGCG GNTCGCAGCT GCGACTCGGC GGGTTGACGG
 551 CAGCGTTGAT GCAGNTCTCG GTACTGGTGC TGCTGCTTTC AGAAATTGGA
 601 AGATAA

5 This encodes a protein having amino acid sequence <SEQ ID 100>:

1 MILLHLDFLS ALLYAAVFLF LIFRAGMLQW FWASIMLWLG ISVLGAKLMP
 51 GIWGMTRAAP LFIPHFYLT LGSIFFFIGHW NRKTDGNGWQ ADPEHPLLGL
 101 FAVSNVSM TLAFVGCALVH YCFSXTVQVF VFAALLKLYA LKPVYWFVLQ
 151 FVLMAYAYVH RCGIDRQPPS TFGGSQLRLG GLTAALMQXS VLVLLLSEIG
 201 R*

ORF18a and ORF18-1 show 99.0% identity in 201 aa overlap:

		10	20	30	40	50	60
15	orf18a.pep	MILLHLDFLSALLYAAVFLFLIFRAGMLQWFWASIMLWLGISVLGAKLMPGIWGMTRAAP					
	orf18-1	MILLHLDFLSALLYAAVFLFLIFRAGMLQWFWASIMLWLGISVLGAKLMPGIWGMTRAAP					
		10	20	30	40	50	60
20	orf18a.pep	LFIPHFYLT LGSIFFFIGHWNRKTDGNGWQADPEHPLLGLFAVSNVSM TLAFVGCALVH					
	orf18-1	LFIPHFYLT LGSIFFFIGHWNRKTDGNGWQADPEHPLLGLFAVSNVSM TLAFVGCALVH					
		70	80	90	100	110	120
25	orf18a.pep	YCFSXTVQVFVFAALLKLYALKPVYWFVLQFVLMAYAYVHRCGIDRQPPSTFGGSQLRLG					
	orf18-1	YCFSXTVQVFVFAALLKLYALKPVYWFVLQFVLMAYAYVHRCGIDRQPPSTFGGSQLRLG					
		130	140	150	160	170	180
30	orf18a.pep	GLTAALMQXS VLVLLLSEIGRX					
	orf18-1	GLTAALMQXS VLVLLLSEIGRX					
35		190	200				

Homology with a predicted ORF from *N.gonorrhoeae*

ORF18 shows 93.1% identity over a 116aa overlap with a predicted ORF (ORF18.ng) from *N. gonorrhoeae*:

40	orf18.pep	GNGWQADPEHPLLGLFAVSNVSM TLAFVGI	30
	orf18ng	TRAAPLFIPHFYLT LGSIFFFIGHWNRKTDGNGWQADPEHPLLGLFAVSNVSM TLAFVGI	115
	orf18.pep	CALVHYCFSGTVQVFVFAALLKLYALKPVYWFVLQFVLMAYAYVHRCGIDRQPPSTFGGS	90
45	orf18ng	CALVHYCFSGTVQVFVFAALLKLYALKPVYWFVLQFVLMAYAYVHRCGIDRQPPSTFGGS	175
	orf18.pep	QLRLGGLTAALMQXS VLVLLLSEIGR	116
	orf18ng	QLRLGVLAALMQVAVTAMLLAEIGR	201

50 The complete length ORF18ng nucleotide sequence is <SEQ ID 101>:

1 ATGATTTTGC TGCATTTGGA TTTTGTGTCT GCCTTACTGt aTGCGGcggt
 51 tttTctgTTT CTGATATTCC GCGCAGGAAT GTTGCAATGG TTTTGGGCGA
 101 GTATTGCGTT GTGGCTCGGC ATCTCGGTTT TAGGGGTAAA GCTGATGCCG
 151 GGGATGTGGG GAATGACCCG CGCCGCGCCT TTGTTTCATCC CCCATTTTAA
 201 CCTGACTTTG GGCAGCATAT TTTTTCAT CCGGTATTGG AACCGGAAAA
 251 CAGATGGAAA CGGATGGCAG GCAGACCCG AACATCCGCT GCTCGGGCTT
 301 TTTGCCGTCA GTAATGTATC GATGACGCTT GCTTTTGTGC GAATATGTGC
 351 GTTGGTG CAT TATTGCTTTT CGGGAACGGT TCAAGTGTTC GTGTTGCGG
 401 CATGCTCAA ACTTTATGCG CTGAAGCCGG TTTATTGGTT CGTGTGTCAG
 451 TTTGTATTGA TGGCGGttgC CTATGTCCAC CGCTGCGGTA TAGACCGGCA
 501 GCCGCCGTCA ACGTTCGGCG GTTCGCAGCT GCGACTCGGC GTGTGCGCG

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551 CGATGTTGAT GCAGGTTGCG GTAACGGCGA TGCTGCTTGC CGAAATCGGC
601 AGATGA

This encodes a protein having amino acid sequence <SEQ ID 102>:

5 1 MILLHLDFLS ALLYAAVFLF LIFRAGMLQW FWASIALWLG ISVLGVKLMP
 51 GMWGMTRAAP LFIPHFYLT L GSIFFFIGYW NRKTDGNGWQ ADPEHPLLGL
 101 FAVSNVSMTL AFVGICALVH YCFSGTVQVF VFAALLKLYA LKPVYWFVLQ
 151 FVLMAYAYVH RCGIDRQPPS TFGGSQRLRG VLAAMLMOVA VTAMLLAEIG
 201 R*

This ORF18ng protein sequence shows 94.0% identity in 201 aa overlap with ORF18-1:

10		10	20	30	40	50	60
	orf18-1.pep	MILLHLDFLSALLYAAVFLFLIFRAGMLQWFWASIMLWLGISVLGAKLMPGIWGMTRAAP					
	orf18ng						
		10	20	30	40	50	60
15		70	80	90	100	110	120
	orf18-1.pep	LFIPHFYLT LGSIFFFIGHWNRKTDGNGWQADPEHPLLGLFAVSNVSMTLAFVGICALVH					
	orf18ng						
20		70	80	90	100	110	120
		130	140	150	160	170	180
	orf18-1.pep	YCFSGTVQVVFVFAALLKLYALKPVYWFVLQFVLMAYAYVHRCGIDRQPPSTFGGSQRLRG					
25	orf18ng						
		130	140	150	160	170	180
		190	200				
	orf18-1.pep	GLTAALMQVSVLVLLSEIGRX					
30	orf18ng	:					
		190	200				

Based on this analysis, including the presence of several putative transmembrane domains in the
35 gonococcal protein, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and
their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 13

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 103>:

40 1 ATGAAAACCC CACTCCTCAA GCCTCTGCTN ATTACCTCGC TTCCCGTTTT
 51 CGCCAGTGTT TTTACCGCCG CCTCCATCGT CTGGCAGCTA GGCGAACCCA
 101 AGCTCGCCAT GCCTTCGTA CTCGGCATCA TCGCCGGCGG CCTGTGCGAT
 151 TTGGACAACC NCNTGACCGG ACGGCTNAAA AACATCATCA CCACCGTCGC
 201 CCTGTTCACC CTCTCCTCGC TCACGGCACA AAGCACCTC GGCACAGGGC
 251 TGCCCTTCAT CCTCGCCATG ACCCTGATGA CTT.CG.CTT CACCATTTTA
45 301 GGCGCGNCG ...

This corresponds to the amino acid sequence <SEQ ID 104; ORF19>:

1 MKTPLLKPLL ITSLPVFASV FTAASIVWL GEPLAMPFV LGIIAGGLVD
51 LDNXXTGRLK NIITTVALEF LSSLTAQSTL GTGLPFILAM TLMTXXTIL
101 GAX...

50 Further work revealed the complete nucleotide sequence <SEQ ID 105>:

1 ATGAAAACCC CACTCCTCAA GCCTCTGCTC ATTACCTCGC TTCCCGTTTT
51 CGCCAGTGTT TTTACCGCCG CCTCCATCGT CTGGCAGCTA GGCGAACCCA
101 AGCTCGCCAT GCCTTCGTA CTCGGCATCA TCGCCGGCGG CCTGTGCGAT
151 TTGGACAACC GCCTGACCGG ACGGCTGAAA AACATCATCA CCACCGTCGC

201	CCTGTTCAACC	CTCTCCTCGC	TCACGGCACA	AAGCACCCCTC	GGCACAGGGC
251	TGCCCTTCAT	CCTCGCCATG	ACCCTGATGA	CCTTCGGCTT	CACCATTTTA
301	GGCGCGGTCG	GGCTCAAATA	CCGCACCTTC	GCCTTCGGTG	CACTCGCCGT
351	CGCCACCTAC	ACCACACTTA	CCTACACCCC	CGAAACCTAC	TGGCTGACCA
401	ACCCCTTCAT	GATTTTATGC	GGCACCGTAC	TGTACAGCAC	CGCCATCCTC
451	CTGTTCCAAA	TCGTCTCGCC	CCACCGCCCC	GTCCAAGAAA	GCGTCGCCAA
501	CGCCTACGAC	GCACTCGGCG	GCTACCTCGA	AGCCAAAGCC	GACTTCTTCG
551	ACCCCGATGA	GGCAGCCTGG	ATAGGCAACC	GCCACATCGA	CCTCGCCATG
601	AGCAACACCG	GCGTCATCAC	CGCCTCAAC	CAATGCCGTT	CCGCCCTGTT
651	TTACCGCCTT	CGCGGCAAAC	ACCGCCACCC	GCGCACCGCC	AAAATGCTGC
701	GTTACTACTT	TGCCGCCCAA	GACATACACG	AACGCATCAG	CTCCGCCAC
751	GTCGATTATC	AGGAAATGTC	CGAAAAATTC	AAAAACACCG	ACATCATCTT
801	CGCATCCAC	CGCCTGCTCG	AAATGCAGGG	ACAAGCCTGC	CGCAACACCG
851	CCCAAGCCCT	GCGCGCAAGC	AAAGACTACG	TTTACAGCAA	ACGCCTCGGC
901	CGCGCCATCG	AAGGCTGCCG	CCAATCGCTG	CGCCTCCTTT	CAGACAGCAA
951	CGACAGTCCC	GACATCCGCC	ACCTGCGCCG	CCTTCTCGAC	AACCTCGGCA
1001	GCGTCGACCA	GCAGTTCGCG	CAACTCCAGC	ACAACGGCCT	GCAGGCAGAA
1051	AACGACCGCA	TGGGCGACAC	CGCATCGCC	GCCCTCGAAA	CCAGCAGCCT
1101	CAAAAACACC	TGGCAGGCAA	TCCGTCCGCA	GCTAAACCTC	GAATCAGGCG
1151	TATTCCGCCA	TGCCGTCCGC	CTGTCCCTCG	TGTTTGCCGC	CGCCTGCACC
1201	ATCGTCGAAG	CCCTCAACCT	CAACCTCGGC	TACTGGATAC	TACTGACCGC
1251	CCTTTTCGTC	TGCCAACCCA	ACTACACCGC	CACCAAAGC	CGCGTCGGCC
1301	AGCGCATCGC	CGGCACCGTA	CTCGGCGTAA	TCGTGCGCTC	GCTCGTCCCC
1351	TACTTCACCC	CGTCTGTCGA	AACCAAACCT	TGGATTGTCA	TGCGCAGTAC
1401	CACCCTCTTT	TTCATGACCC	GCACCTACAA	ATACAGTTTC	TCCACCTTCT
1451	TCATTACCAT	TCAAGCCTCG	ACCAGCCTCT	CCCTCGCAGG	TTTGGACGTA
1501	TACGCCGCCA	TGCCCGTAGC	CATCATCGAC	ACCATTATCG	GCGCATCCCT
1551	TGCCTGGGCG	GCAGTCAGCT	ACCTGTGGCC	AGACTGGAAA	TACCTCACGC
1601	TCGAACGCAC	CGCCGCCCTT	GCCGTATGCA	GCAACGGTGC	CTATCTCGAA
1651	AAAATCACCG	AACGCCTCAA	AAGCGGCGAA	ACCGGCGACG	ACGTGCAATA
1701	CCGCGCCACC	CGCCGCCGCG	CCCACGAACA	CACCGCCGCC	CTCAGCAGCA
1751	CCCTTTCCGA	CATGAGCAGC	GAACCCGCAA	AATTGCGCCA	CAGCCTGCAA
1801	CCCGGCTTTA	CCCTGCTCAA	AACCGGCTAC	GCCCTGACCG	GCTACATCTC
1851	CGCCCTCGGC	GCATACCGCA	GCGAAATGCA	CGAAGATGCG	AGCCCCGACT
1901	TTACCGCACA	GTTCCACCTC	GCCGCCGAAC	ACACCGCCCA	CATCTTCCAA
1951	CGFTLLKTY	ALTGYISALG	AYRSEMHEEC	SPDFTAQFHL	AAEHTAHIFQ
2001	GCGCGGCGAA	CTCGACACCC	TCCGCACCCA	CAGCAGCGGA	ACACAAAGCC
2051	ACATCCTCCT	CCAACAGCTC	CAACTCATCG	CCCGACAGCT	CGAACCCTAC
2101	TACCGCGCCT	ACCGCCAAAT	TCCGCACAGG	CAGCCCCAAA	ATGCAGCCTG
2151	A				

This corresponds to the amino acid sequence <SEQ ID 106; ORF19-1>:

1	MKTPLLKPLL	ITSLPVFASV	FTAASIVWQL	GEPKLAMPFV	LGIIAGGLVD
51	LDNRLTGRLL	NIITTVALFT	LSSLTAQSTL	GTGLPFILAM	TLMTFGFTIL
101	GAVGLKYRTF	AFGALAVATY	TTLTYTPETY	WLTNPFMILC	GTVLYSTAIL
151	LFQIVLPHRP	VQESVANAYD	ALGGYLEAKA	DDFDPDEAAW	IGNRRHIDLAM
201	SNTGVITAFN	QCRSALFYRL	RGKHRHERTA	KMLRYYFAAQ	DIHERISSAH
251	VDYQEMSEKF	KNTDIIFRIH	RLLEMOGOAC	RNTAQALRAS	KDYVYSKRLG
301	RAIEGCRQSL	RLSDSNDSP	DIRHLRRLLD	NLGSVDQOFR	QLQHNGLQAE
351	NDRMGDTRIA	ALETSSLKNT	WQAIRPOLNL	ESGVFRHAVR	LSLVVAAACT
401	IVEALNLLNG	YWILLTALFV	QOPNYTATKS	RVRQRIAGTV	LGVIVGSLVP
451	YFTPSVETKL	WIVIASTTLF	FMTRTYKYSF	STFFITIQAL	TSLSLAGLDV
501	YAAMPVRIID	TIIGASLAWA	AVSYLWPDWK	YLTLERTAAL	AVCSNGAYLE
551	KITERLKSGE	TGDDVEYRAT	RRRAHEHTAA	LSSTLSDMSS	EPAKFADSLQ
601	PGFTLLKTY	ALTGYISALG	AYRSEMHEEC	SPDFTAQFHL	AAEHTAHIFQ
651	HLPETEPDDF	QTALDTLRGE	LDTLRTHSSG	TQSHILLQQL	QLIARQLEPY
701	YRAYRQIPHR	QPQNAA*			

Computer analysis of this amino acid sequence gave the following results:

Homology with predicted transmembrane protein YHFK of *H. influenzae* (accession number P44289)

ORF19 and YHFK proteins show 45% aa identity in 97 aa overlap:

60	orf19	6	LKPLLITSLPVFASVFTAASIVWQLGEPKLAMPFVLGIIAGGLVDLDNXXTGRLKNIITT	65
			L +I+++PVF +V AA +W +MP +LGIIAGGLVDLDN TGRLKN+ T	
	YHFK	5	LNAKVISTIPVFIAVNIAAVGIWFFDISQSMLILGIIAGGLVDLDNRLTGRLKNVFFT	64

ORF19 shows 92.2% identity over a 102aa overlap with an ORF (ORF19a) from strain A of *N.*

		10	20	30	40	50	60
10	orf19.pep	<u>MKTPLLKPLLITSLPVFASVFTAASIVWQLGEPKLA</u> <u>MPFVLGI</u> <u>IAGGLVDL</u> <u>DNXXTGRLK</u>					
	orf19a	<u>MKTPPCLKPLLITSLPVFASVFTAASIVWQLGEPKLA</u> <u>MPFVLGI</u> <u>IAGGLVDL</u> <u>DNRLTGRLK</u>					
		10	20	30	40	50	60
15		70	80	90	100		
	orf19.pep	<u>NIITTVALFTLSSSLTAQSTLGTGLPFI</u> <u>LAMTLM</u> <u>TXXTILGAX</u>					
		:	:	:	:	:	:
	orf19a	<u>NIIATVALFTLSSSLVAQSTLGTGLPFI</u> <u>LAMTLM</u> <u>TFGTIMGAV</u> <u>GLKYRTFAFGALAVATY</u>					
		70	80	90	100	110	120
20	orf19a	<u>TTLTYTPETYWLTNPF</u> <u>MLICGTVLYSTAIL</u> <u>EQIILPHRPVQEN</u> <u>VANAYEALGSYLEAKA</u>					
		130	140	150	160	170	180

	1	ATGAAAACCC	CACCCCTCAA	GCCTCTGCTC	ATTACCTCGC	TTCCCGTTTT
25	51	CGCCAGTGTC	TTTACGCGCG	CCTCCATCGT	CTGGCAGCTG	GGCGAACCCA
	101	AGCTCGCCAT	GCCCTTCGTA	CTCGGCATCA	TCGCTGGCGG	CCTGGTTCGAT
	151	TTGGACAACC	GCCGTGACCG	ACGGCTGAAA	AACATCATCG	CCACCGTCGC
	201	CCTGTTACAC	CTCTCCTCAC	TTGTGCGGCA	AAGCACCCCTC	GGCACAGGTT
30	251	TGCCATTCAT	CCTCGCCATG	ACCCTGATGA	CTTTTCGGCTT	TACCATCATG
	301	GGCGCGGTCG	GGCTGAAATA	CCGCACCTTC	GCCTTCGGCG	CATCTCGCGT
	351	CGCCACCTAC	ACCACACTTA	CCTACACCCC	CGAAACCTAC	TGGCTGACCC
	401	ACCCCTTTAT	GATTCTGTGC	GGAACCGTAC	TGTACAGCAC	CGCCATCATC
35	451	CTGTTCCAAA	TCATCCTGCC	CCACCGCCCC	GTTCAAGAAA	AGCTCGCCAA
	501	CGCCTACGAA	GCATCTGGCA	GCTACCTCGA	AGCCAAAGCC	CACTTTTTCG
	551	ATCCCGACGA	AGCCGAATGG	ATAGGCAACC	GCCACATCGA	CCTCGCCATG
	601	AGCAACACCG	GCGTCATCAC	CGCCTTCAAC	CAATGCCGTT	CCGCCCTGTT
40	651	TTACCGCCTT	CGCGGCAAA	ACCGCCACCC	GCGCACCGCC	AAAATGCTGC
	701	GCTACTACTT	CGCGGCCCAA	GACATACACG	AACGCATCAG	CTCCGCCACG
	751	GTCGACTACC	AAGAGATGTC	CGAAAAATTC	AAAAACACCG	ACATCATCTT
	801	CGCATCCAC	CGCCTGCTCG	AAATGCAGGG	ACAAGCCTGC	CGCAACACCG
45	851	CCCAAGCCCT	CGCGCAAGC	AAAGACTACG	TTTACAGCAA	ACGCCCTCGC
	901	CGCGCCATCG	AAGGCTCGCG	CCAATCGCTG	CGCCTCCTTT	CAGACAGCAA
	951	CGACAATCCC	GACATCCGCC	ACCTGCGCGG	CCTTCTCGAC	AACCTCGGCA
	1001	GCGTCGACCA	GCAATTCCCG	CAACTCCAGC	ACAACGGCCT	CGAGGCAGAA
50	1051	AACGACCGCA	TGGGCGACAC	CCGCATCGCC	CGCCTCGAAA	CCGCGACGCT
	1101	CAAAAACACC	TGGCAGGCAG	TCCGTCCGCA	GCTAAACCTC	GAATCAGGCG
	1151	TATTCCGCCA	TGCCGTCCGC	CTGTCCCTTG	TCGTTTGGCG	CGCCTGCACC
	1201	ATCGCTGAAG	CCCTCAACCT	CAACCTCGGC	TACTGGATAC	TACTGACCGC
55	1251	CCTTTTCGTC	TGCCAACCCA	ACTACACCGC	CACCAAAGCG	CGCGTCCGCC
	1301	AGCGCATCGC	CGGCACCGTA	CTCGGCGTAA	TCGTGCGCTC	GCTCGTCCCC
	1351	TACTTTACCC	CCTCCGTGCA	AACCAAACCT	TGGATCGTCA	TCGCCAGTAC
	1401	CACCCCTCTT	TTCATGACCC	GCACCTACAA	ATACAGCTTC	TCGACATTTT
60	1451	TCATACCCAT	TCAAGCCCTG	ACCAGCCTCT	CCCTCGCAGG	GTTGGACGTA
	1501	TACGCGGCCA	TGCCCGTACG	CATCATCGAC	ACCATTATCG	GCGCATCCCT
	1551	TGCTCTGGGCG	GCAGTCAGCT	ACCTGTGGCC	AGACTGGAAA	TACCTCACGC
	1601	TCGAACGCAC	CGCGCCCTTT	GCGCTATGCA	GCAACGCGGC	CTATCTCGAA
65	1651	AAAATCACCG	AACGCCCTCA	AAGCGGCGAA	ACCGGCGACG	ACGTCAATAA
	1701	CGCGGCCACC	CGCGGCCGCG	CCCACGAACA	CACCGCCGCG	CTCAGACGCA
	1751	CCCTTTCGCA	CATGAGCAGC	GAACCCGCAA	AATTTCGCCA	CAGCCTGCAA
	1801	CCCGGCTTTA	CCCTGCTCAA	AACCGGCTAC	GCGCTGACCG	GCTACATCTC
70	1851	CGCCCTCGGC	GCATACCGCA	GCGAAATGCA	CGAAGAATGC	AGCCCCGACT
	1901	TTACCGCACA	GTTCCACCTC	GCGCGCGAAG	ACACCGCCCA	CATCTTCCAA
	1951	CACCTGCCCC	AAACCGAACC	CGACGACTTT	CAGACAGCAC	TGGATACTAT
	2001	GCGCGGCGAA	CTCGACACCC	TCCGCACCCA	CAGCAGCGGA	ACACAAAGCC
75	2051	ACATCCTCCT	CCAACAGCTC	CAACTCATCG	CCCGGCAGCT	CGAACCCTAC
	2101	TACCGCGCCT	ACCGACAAAT	TCCGCACAGG	CAGCCCCAAA	ACGCAGCCTG
	2151	A				

This encodes a protein having amino acid sequence <SEQ ID 108>:

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      1  MKTPPLKPLL  ITSLPVFASV  FTAASIVWQL  GEPKLAMPFV  LGIIAGGLVD
    51  LDNRLTGRLE  NIIATVALFT  LSSLVAQSTL  GTCLPFIAM  TLMTFGFTIM
   101  GAVGLKYRTF  AFGALAVATY  TTLTYTPETY  WLTNPFMILC  GTVLYSTAI
    5  151  LFQIILPHRP  VQENVANAYE  ALGSYLEAKA  DFFDPDEAEW  IGNRHIDLAM
   201  SNTGVITAFN  QCRSALFYRL  RGKHRHPRTA  KMLRYFFAAQ  DIHERISSAH
   251  VDYQEMSEKF  KNTDIIFRIH  RLLEMQQQAC  RNTAQALRAS  KDYVYSKRLG
   301  RAIEGCRQSL  RLLSDSNDNP  DIRHLRRLLD  NLGSVDQQFR  QLQHNLQAE
   351  NDRMGDTRIA  ALETGSLKNT  WQAIRPOLNL  ESGVFRHAVR  LSLVVAAACT
   401  IVEALNLNLG  YWILLTALFV  CQPNYTATKS  RVRQRIAGTV  LGVIVGSLVP
  10  451  YETPSVETKL  WIVIASTTLF  FMTRTYKYSF  STFFITIQAL  TSLSLAGLDV
   501  YAAMPVRIID  TIIGASLAWA  AVSYLWPDWK  YLTLETAAL  AVCSNGAYLE
   551  KITERLKSGE  TGDDVEYRAT  RRRRAHEHTAA  LSSTLSDMSS  EPAKFADSLQ
   601  PGFTLLKGTG  ALTGYISALG  AYRSEMHEEC  SPDFTAQFHL  AAHTAHIFQ
  15  651  HLPETEPDDF  QTALDTLRGE  LDTLRTHSSG  TQSHILLQQL  QLIARQLEPY
   701  YRAYRQIPHR  QPQNAA*

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ORF19a and ORF19-1 show 98.3% identity in 716 aa overlap:

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      10      20      30      40      50      60
 20  orf19a.pep  MKTPPLKPLLITSLPVFASVFTAASIVWQLGEPKLAMPFVLGIIAGGLVDLDNRLTGRLE
      10      20      30      40      50      60
      orf19-1  MKTPPLKPLLITSLPVFASVFTAASIVWQLGEPKLAMPFVLGIIAGGLVDLDNRLTGRLE

      70      80      90     100     110     120
 25  orf19a.pep  NIIATVALFTLSSLVAQSTLGTGLPFIAMTLMTFGFTIMGAVGLKYRTFAFGALAVATY
      70      80      90     100     110     120
      orf19-1  NIITVALFTLSSLTAQSTLGTGLPFIAMTLMTFGFTILGAVGLKYRTFAFGALAVATY

      130     140     150     160     170     180
 30  orf19a.pep  TTLTYTPETYWLTNPFMILCGTVLYSTAILLFQIILPHRPVQENVANAYEALGSYLEAKA
      130     140     150     160     170     180
      orf19-1  TTLTYTPETYWLTNPFMILCGTVLYSTAILLFQIVLPHRPVQESVANAYDALGGYLEAKA

      190     200     210     220     230     240
 35  orf19a.pep  DFFDPDEAEWIGNRHIDLAMSNITGVITAFNQCRSALFYRLRGKHRHPRTAKMLRYFFAAQ
      190     200     210     220     230     240
      orf19-1  DFFDPDEAAWIGNRHIDLAMSNITGVITAFNQCRSALFYRLRGKHRHPRTAKMLRYFFAAQ

      250     260     270     280     290     300
 40  orf19a.pep  DIHERISSAHVDYQEMSEKFKNTDIIIFRIHRLLEMQQQACRNTAQALRASKDYVYSKRLG
      250     260     270     280     290     300
      orf19-1  DIHERISSAHVDYQEMSEKFKNTDIIIFRIHRLLEMQQQACRNTAQALRASKDYVYSKRLG

      310     320     330     340     350     360
 50  orf19a.pep  RAIEGCRQSLRLLSDSNDNPDIRHLRRLLDNLGSVDQQFRQLQHNLQAEENDRMGDTRIA
      310     320     330     340     350     360
      orf19-1  RAIEGCRQSLRLLSDSNDSPDIRHLRRLLDNLGSVDQQFRQLQHNLQAEENDRMGDTRIA

      370     380     390     400     410     420
 55  orf19a.pep  ALETGSLKNTWQAIRPOLNLESGVFRHAVRLSLVVAAACTIVEALNLNLGYWILLTALFV
      370     380     390     400     410     420
      orf19-1  ALETSSLKNTWQAIRPOLNLESGVFRHAVRLSLVVAAACTIVEALNLNLGYWILLTALFV

      430     440     450     460     470     480
 60  orf19a.pep  CQPNYTATKSRVRQRIAGTVLGIVGSLVPYFTPSVETKLWIVIASTTLFEMTRTYKYSF
      430     440     450     460     470     480
      orf19-1  CQPNYTATKSRVRQRIAGTVLGIVGSLVPYFTPSVETKLWIVIASTTLFEMTRTYKYSF

      490     500     510     520     530     540
 65  orf19a.pep  STFFITIQALTSLSLAGLDVYAAMPVRIIDTIIGASLAWAAVSYLWPDWKYLTLETAAL
      490     500     510     520     530     540

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	orf19-1	STFFITIQA	LTSLAGLDV	YAAMPVRI	IDTIIGASL	AWAAVSYL	WPDWKYL	TLERTAAL
		490	500	510	520	530	540	
5	orf19a.pep	AVCSNGAYLEK	ITERLKSGET	GDDVEYRATR	RRRAHEHTA	AALSSTLS	SDMSSEPA	KFADSLO
	orf19-1	AVCSNGAYLEK	ITERLKSGET	GDDVEYRATR	RRRAHEHTA	AALSSTLS	SDMSSEPA	KFADSLO
10	orf19a.pep	PGFTLLKTGY	ALTGYISAL	GAYRSEMHEE	CSPDFTAQ	FHLAAEHT	AHIFQHL	PETEPDDF
	orf19-1	PGFTLLKTGY	ALTGYISAL	GAYRSEMHEE	CSPDFTAQ	FHLAAEHT	AHIFQHL	PETEPDDF
15	orf19a.pep	QTALDTLRG	ELDTLRTH	SSGTSQSH	ILLQQLQ	LQLIARQ	LEPYRAYR	QIPHRRQ
	orf19-1	QTALDTLRG	ELDTLRTH	SSGTSQSH	ILLQQLQ	LQLIARQ	LEPYRAYR	QIPHRRQ
20	orf19a.pep	QALDTLRG	ELDTLRTH	SSGTSQSH	ILLQQLQ	LQLIARQ	LEPYRAYR	QIPHRRQ
	orf19-1	QALDTLRG	ELDTLRTH	SSGTSQSH	ILLQQLQ	LQLIARQ	LEPYRAYR	QIPHRRQ

Homology with a predicted ORF from *N.gonorrhoeae*

ORF19 shows 95.1% identity over a 102aa overlap with a predicted ORF (ORF19.ng) from *N. gonorrhoeae*:

25	orf19.pep	MKTPLLKPLL	ITSLPVFASV	FTAASIVWQL	GEPKlampFV	LGIAGGLVD	LDNXXTGR	LK	60
	orf19ng	MKTPLLKPLL	ITSLPVFASV	FTAASIVWQL	GEPKlampFV	LGIAGGLVD	LDNRLTGR	LK	60
30	orf19.pep	NIITVAF	FTLSSSLTAQ	STLGTGLP	FILAMT	LMTXXFT	ILGAX		103
	orf19ng	NIIATVA	FTLSSSLTAQ	STLGTGLP	FILAMT	LMTFGFT	ILGAVGL	KYRTFA	FGALAVATY 120

An ORF19ng nucleotide sequence <SEQ ID 109> is predicted to encode a protein having amino acid sequence <SEQ ID 110>:

35	1	MKTPLLKPLL	ITSLPVFASV	FTAASIVWQL	GEPKlampFV	LGIAGGLVD	
	51	LDNRLTGR	LK NIIATVA	FTLSSSLTAQ	STLGTGLP	FILAMT	LMTFGFT
40	101	GAVGLKYR	TF AFGALAV	ATY TLT	YTPETY	WLTNP	FILC GTVLY
	151	LFQIILPH	RP VQESVAN	AYE ALGGY	LEAKA DFFD	PDEAAW	IGNRH
40	201	NTGVITAF	N QCRSAL	FYRL R	GKHRH	PRTA K	MLRYF
	251	VDYQEMSE	KF KNTDII	FRIR R	LLEMQ	GQAC R	NTAQAI
40	301	RAIEGCRQ	SL RLLSD	GNDSP	DIRHLS	RLLD NLG	SVDQ
	351	NDRMGD	TRIA A	LETGS	FKNT *		

Further work revealed the complete nucleotide sequence <SEQ ID 111>:

45	1	ATGAAAACCC	CACTCCTCAA	GCCTCTGCTC	ATTACCTCGC	TTCCCGTTTT	
	51	CGCCAGTGTC	TTTACCGCCG	CCTCCATCGT	CTGGCAGCTA	GGCGAACCCA	
50	101	AGCTCGCCAT	GCCCTTCGTA	CTCGGCATCA	TCGCCGGCGG	CCTGGTCGAT	
	151	TTGGACAACC	GCCTGACCGG	ACGGCTGAAA	AACATCATCG	CCACCGTCGC	
50	201	CCTGTTTACC	CTCTCCTCGC	TCACGGCGCA	AAGCACCTC	GGCACAGGGC	
	251	TGCCCTTCAT	CCTCGCCATG	ACCCTGATGA	CCTTCGGCTT	TACCATTTTA	
50	301	GGCGCGGTCG	GGCTGAAATA	CCGCACCTTC	GCCTTCGGCG	CACTCGCCGT	
	351	CGCCACCTAC	ACCACGCTTA	CCTACACCCC	CGAAACCTAC	TGGCTGACCA	
55	401	ACCCCTTCAT	GATTTTATGC	GGCACCGTAC	TGTACAGCAC	CGCCATCATC	
	451	CTGTTCCAAA	TCATCCTGCC	CCACCGCCCC	GTCCAAGAAA	GGCTCGCCAA	
55	501	TGCTACGAA	GCACTCGGCG	GCTACCTCGA	AGCCAAAGCC	GACTTCTTCG	
	551	ACCCCGATGA	GGCAGCCTGG	ATAGGCAACC	GCCACATCGA	CCTCGCCATG	
60	601	AGCAACACCG	GCGTCATCAC	CGCCTTCAAC	CAATGCCGTT	CCGCCCTGTT	
	651	TTACCGTTTG	CGCGGCAAAC	ACCGCCACCC	GCGCACCGCC	AAAATGCTGC	
60	701	GCTACTACTT	CGCCGCCCAA	GACATCCACG	AACGCATCAG	CTCCGCCAC	
	751	GTCGACTACC	AAGAGATGTC	CGAAAAATTC	AAAAACACCG	ACATCATCTT	
60	801	CCGCATCCGC	CGCCTGCTCG	AAATGCAGGG	GCAGGCGTGC	CGCAACACCG	
	851	CCCAAGCCAT	CCGTCGGGCG	AAAGACTAcg	tTTACAGCAA	ACGCCCTCGGA	
60	901	CGGCCATcg	aaggctgCGG	CCAGTCGCTg	cgcctCCTTt	cagacggcaA	
	951	CGACAGTCCC	GACATCCGCC	ACCTGAGcgcg	CCTTCTCGAC	AACCTCGgca	

1001 GCGTcgacca gcagtTCgc caactCCGAC ACAGcgactC CCCCCcgaa
 1051 Aacgaccgca tggcgacac CCGCATCGCC GCCCtcgaaa cggcgagctT
 1101 caaaaaCAcc tggcaggCAA TCCGTCCGCa gctgaaCCTC GAATCatgCG
 1151 TATTCGGCCA TGCCGTCCGC CTGTCCCTCG TCGTTGCCGC CGCCTGCACC
 1201 ATCGTCgaag cCCTCAACCT CAACCTCGGC TACTGGATAC TGCTGACCGC
 1251 CCTTTTCGTC TGCCAACCCA ACTACACCGC CACCAAAAGC CGCGTGACC
 1301 AACGCATCGC CGGCACCGTA CTCGGCGTAA TCGTCGGGTC GTCGTCCCC
 1351 TACTTCACCC CCTCCGTCTGA AACCAAATC TGGATTGTCA TCGCCGGTAC
 1401 CACCTGTTC TTCATGACCC GCACCTACAA ATACAGTTTC TCCACCTTCT
 1451 TCATCACCAT TCAGGCACTG ACCAGCCTCT CCCTCGCAGG TTTGGACGTA
 1501 TACGCCGCCA TGCCCGTGCG CATCATcgac ACCATTATCG GCGCATCCCT
 1551 TGCCTGGGCG GCGGTGAGCT ACCTGTGGCC AGACTGGAAA TACCTCAGC
 1601 TCGAACGCAC CGCCGCCCTT GCCGTATGCA GCAGCGGCAC ATACCTCCAA
 1651 AAAATTGCCG AACGCCTCAA AACGGCGGAA ACCGGCGACG ACATAGAATA
 1701 CCGCATCACC CGCCGCCGCG CCCACGAACA CACCGCGGCC CTCAGCAGCA
 1751 CCCTTTCGGA CATGAGCAGC GAACCGCAA AATTGCGCGA CAGCTGCAA
 1801 CCGGCTTTA CCTGTCTCAA AACCGGCTAC GCCCTGACCG GCTACATCTC
 1851 CGCCCTCGGC GCATACCGCA GCGAAATGCA CGAAGAATGC AGCCCCGACT
 1901 TTACCGCACA GTTCCACCTT GCCGCCGAAC ACACCGCCCA CATCTTCCAA
 1951 CACCTGCCCG ACATGGGACC CGACGACTTT CAGACGGCAT TGGATACACT
 2001 GCGCGGCGAA CTCGGCACCC TCCGCACCCG CAGCAGCGGA ACACAAAGCC
 2051 ACATCCTCCT CCAACAGCTC CAACTCATCG CccgGCAACT CGAACCTTAC
 2101 TACCGCGCCT ACCGACAAAT TCCGCACAGG CAGCCCCAAA ACGCAGCCTG
 2151 A

25 This corresponds to the amino acid sequence <SEQ ID 112; ORF19ng-1>:

1 MKTPLLKPLL ITSLPVFASV FTAASIVWQL GEPKlampfv LGIIAGGLVD
 51 LDNRLTGRlk NIIATVALFT LSSLTAQSTL GTGLPFILAM TLMTFGFTIL
 101 GAVGLKYRTF AFGALAVATY TTLTYTPETY WLTNPFMILC GTVLYSTAIL
 151 LFQIILPHRP VQESVANAYE ALGGYLEAKA DFFDPDEAAW IGNRHIDLAM
 201 SNTGVITAFN QCRSALFYRL RGKHRHPRTA KMLRYFFAAQ DIHERISSAH
 251 VDYQEMSEKF KNTDIIIFRIR RLLEMQGGAC RNTAQAIRSG KDYVYSKRLG
 301 RAIEGCRQSL RLSDGNDSP DIRHLSRLLD NLGSVDQQFR QLRHSDSPA
 351 NDRMGDTRIA ALETGSFKNT WQAIRPQLNL ESCVFRHAVR LSLVVAAC
 401 IVEALNLNLG YWILLTALFV CQPNYATKS RVYQRIAGTV LGVIVGSLVP
 451 YETPSVETKL WIVIAGTTLF FMTRYKYSE STFFITIQL TSLSLAGLDV
 501 YAAMPVRIID TIIIGASLAWA AVSYLWPDWK YLTLERTAAL AVCSSGTYLQ
 551 KIAERLKTGE TGDDIEYRIT RRRRAHEHTAA LSSTLSDMSS EPAKFADSLQ
 601 PGFTLLKTGY ALTGYISALG AYRSEMHEEC SPDFTAQFHL AAEHTAHIFQ
 651 HLPDMGPDDF QTALDTLRGE LGTLRTRSSG TQSHILLQQL QLIARQLEPY
 701 YRAYRQIPHR QPQNAA*

ORF19ng-1 and ORF19-1 show 95.5% identity in 716 aa overlap:

		10	20	30	40	50	60
orf19-1.pep		MKTPLLKPLLITSLPVFASVFTAASIVWQLGEPKlampfvLGIIAGGLVDLDNRLTGRlk					
orf19ng-1		MKTPLLKPLLITSLPVFASVFTAASIVWQLGEPKlampfvLGIIAGGLVDLDNRLTGRlk					
		10	20	30	40	50	60
orf19-1.pep		NIIITVALFTLSSLTAQSTLGTGLPFILAMTLMTFGFTILGAVGLKYRTFAFGALAVATY					
orf19ng-1		NIIATVALFTLSSLTAQSTLGTGLPFILAMTLMTFGFTILGAVGLKYRTFAFGALAVATY					
		70	80	90	100	110	120
orf19-1.pep		TTLTYTPETYWLTNPFMILCGTVLYSTAILLFQIIVLPHRPVQESVANAYDALGGYLEAKA					
orf19ng-1		TTLTYTPETYWLTNPFMILCGTVLYSTAILLFQIILPHRPVQESVANAYEALGGYLEAKA					
		130	140	150	160	170	180
orf19-1.pep		TTLTYTPETYWLTNPFMILCGTVLYSTAILLFQIIVLPHRPVQESVANAYDALGGYLEAKA					
orf19ng-1		TTLTYTPETYWLTNPFMILCGTVLYSTAILLFQIILPHRPVQESVANAYEALGGYLEAKA					
		190	200	210	220	230	240
orf19-1.pep		DFFDPDEAAWIGNRHIDLAMSNITGVITAFNQCRSALFYRLRGKHRHPRTAKMLRYFFAAQ					
orf19ng-1		DFFDPDEAAWIGNRHIDLAMSNITGVITAFNQCRSALFYRLRGKHRHPRTAKMLRYFFAAQ					
		250	260	270	280	290	300
orf19-1.pep		DIHERISSAHVDYQEMSEKFKNITDIIIFRIHRLLEMQGGACRNTAQALRASKDYVYSKRLG					

	orf19ng-1	: : : DIHERISSAHVDYQEMSEKFKNTDIIFRIRRLLEMGGQACRNTAQAIRSGKDYVYSKRLG 250 260 270 280 290 300
5	orf19-1.pep	310 320 330 340 350 360 RAIEGCRQSLRLLSDSNDSPDIRHLRRLLDNLGSVDQQFRQLQHNGLOAENDRMGDTRIA : : : :
10	orf19ng-1	310 320 330 340 350 360 RAIEGCRQSLRLLSDGNDSPDIRHLSRLLDNLGSVDQQFRQLRHSDSPAENDRMGDTRIA
	orf19-1.pep	370 380 390 400 410 420 ALETSSLKNTWQAIROPQLNLESGVFRHAVRLSLVVAACTIVEALNINLGYWILLTALFV : : : :
15	orf19ng-1	370 380 390 400 410 420 ALETGSFKNTWQAIROPQLNLESCVFRHAVRLSLVVAACTIVEALNINLGYWILLTALFV
	orf19-1.pep	430 440 450 460 470 480 CQPNYTATKSRVRQRIAGTVLGVIVGSLVPYFTPSVETKLWIVIASITLFFMTRTYKYSF : : : :
20	orf19ng-1	430 440 450 460 470 480 CQPNYTATKSRVYQRIAGTVLGVIVGSLVPYFTPSVETKLWIVIASITLFFMTRTYKYSF
	orf19-1.pep	490 500 510 520 530 540 STFFITIQAALTSLSLAGLDVYAAMPVRIIDTIIGASLAWAAVSYLWPDWKYLTLETAAL : : : :
25	orf19ng-1	490 500 510 520 530 540 STFFITIQAALTSLSLAGLDVYAAMPVRIIDTIIGASLAWAAVSYLWPDWKYLTLETAAL
	orf19-1.pep	550 560 570 580 590 600 AVCSNGAYLEKITERLKSGETGDDVEYRATRRRAHEHTAALSSTLSDSMSSEPAKFADSLQ : : : :
30	orf19ng-1	550 560 570 580 590 600 AVCSSGTYLQKIAERLKTGETGDDIEYRITRRRAHEHTAALSSTLSDSMSSEPAKFADSLQ
	orf19-1.pep	610 620 630 640 650 660 PGFTLLKTGYALTGYISALGAYRSEMHEECSPDFTAQFHLAAEHTAHIFQHLPETEPDDF : : : :
35	orf19ng-1	610 620 630 640 650 660 PGFTLLKTGYALTGYISALGAYRSEMHEECSPDFTAQFHLAAEHTAHIFQHLPDMGPDDF
	orf19-1.pep	670 680 690 700 710 QTALDTLRGELDTLRTHSSGTQSHILLQQLLIARQLEPYRAYRQIPHRQPQNAAX : : : :
40	orf19ng-1	670 680 690 700 710 QTALDTLRGELGTLRTRSSGTQSHILLQQLLIARQLEPYRAYRQIPHRQPQNAAX
45		670 680 690 700 710

In addition, ORF19ng-1 shows significant homology to a hypothetical gonococcal protein previously entered in the databases:

50	sp O33369 YOR2_NEIGO HYPOTHETICAL 45.5 KD PROTEIN (ORF2) gnl PID e1154438 (AJ002423) hypothetical protein [Neisseria gonorrh] Length = 417 Score = 1512 (705.6 bits), Expect = 5.3e-203, P = 5.3e-203 Identities = 301/326 (92%), Positives = 306/326 (93%)
55	Query: 307 RQSLRLLSDGNDSPDIRHLSRLLDNLGSVDQQFRQLRHSDSPAENDRMGDTRIAALETGS 366 RQSLRLLSDGNDSPDIRHLSRLLDNLGSVDQQFRQLRHSDSPAENDRMGDTRIAALETGS Sbjct: 1 RQSLRLLSDGNDSPDIRHLSRLLDNLGSVDQQFRQLRHSDSPAENDRMGDTRIAALETGS 60
60	Query: 367 FKNTWQAIROPQLNLESCVFRHAVRLSLVVAACTIVEALNINLGYWILLTALFVCQPNYT 426 FKNTWQAIROPQLNLESCVFRHAVRLSLVVAACTIVEALNINLGYWILLTALFVCQPNYT Sbjct: 61 FKNTWQAIROPQLNLESGVFRHAVRLSLVVAACTIVEALNINLGYWILLTALFVCQPNYT 120
65	Query: 427 ATKSRVYQRIAGTVLGVIVGSLVPYFTPSVETKLWIVIASITLFFMTRTYKYSFSTFFIT 486 ATKSRVYQRIAGTVLGVIVGSLVPYFTPSVETKLWIVIASITLFFMTRTYKYSFSTFFIT Sbjct: 121 ATKSRVYQRIAGTVLGVIVGSLVPYFTPSVETKLWIVIASITLFFMTRTYKYSFSTFFIT 180
	Query: 487 IQALTSLSLAGLDVYAAMPVRIIDTIIGASLAWAAVSYLWPDWKYLTLETAALAVCSSG 546 IQALTSLSLAGLDVYAAMPVRIIDTIIGASLAWAAVSYLWPDWKYLTLETAALAVCSSG Sbjct: 181 IQALTSLSLAGLDVYAAMPVRIIDTIIGASLAWAAVSYLWPDWKYLTLETAALAVCSSG 240

Query: 547 TYLQKIAERLKTGETGDDIEYRITRRRAHEHTAALSSSTLSDMSSEPAKFADSLQPGFTLL 606
 TYLQKIAERLKTGETGDDIEYRITRRRAHEHTAALSSSTLSDMSSEPAKFAD+ P
 Sbjct: 241 TYLQKIAERLKTGETGDDIEYRITRRRAHEHTAALSSSTLSDMSSEPAKFADTCNPALPCS 300

5 Query: 607 KTGyALTGYISALGAYRSEMHEECSP 632
 K ALTGYISALG ++ + +P
 Sbjct: 301 KPATALGYISALGHATAAKCTKNAAP 326

Based on this analysis, including the presence of several putative transmembrane domains in the gonococcal protein (the first of which is also seen in the meningococcal protein), and on homology with the YHFK protein, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 14

The following DNA sequence, believed to be complete, was identified in *N.meningitidis* <SEQ ID 113>:

```

15      1  ATGAATATGC TGGGAGCTTT GGCAAAAGTC GGCAGCCTGA CGATGGTGTG
      51  GCGCGTTTTG GGATTGTGTC GCGATACGGT CATTGCGCGG GCATTCGGCG
     101  CGGGTATGGC GACGGATGCG TTTTGTGTCG CGTTCAAACT GCCCAACCTG
     151  CTTGCGCCGC TGTTTGCGGA GGGGGCGTTT GCCCAAGCGT TTGTGCCGAT
     201  TTTGCGCGAA TACAAGGAAA CGCGTTCAAA AGAGGCGG.C GAAGCCTTTA
     251  TCCGCCATGT GCGGGGATG CTGTCGTTTG TACTGGTTAT CGTTACCGCG
     301  CTGGGCATAC TTGCCGCGCC TTGGGTGATT TATGTTCCG CACCCGAGTT
     351  TTGCCCAAGA TGCCGACAAA TTTCAGCTCT CCATCGATTG GCTGCGGATT
     401  ACGTTTCCTT ATATATTATT GATTTCCTCG TCTTCATTG TCGGCTCGGT
     451  ACTCAATTCT TATCATAAGT TCGGCATTCC GCGGTTTACG CCAC.GTTTC
     501  TGAACGTGTC GTTTATCGTA TTCGCGCTGT TTTTCGTGCC GTATTTCGAT
     551  CCGCCCGTTA CCGCGCyGGC GTGGGCGGTC TTTGTGCGCG GCATTTTGCA
     601  ACTCGmTTC CAACTGCCCT GGCTGGCGAA ACTGGGCTTT TTGAACTGCG
     651  CCAAACTGAG TTTCAAAGAT GCGGCGGTCA ACCGCGTGAT GAAACAGATG
     701  GCGCCTGCGA TTTTgGCGT GAAGTGGCG CAGGTTTCTT TGGTGATCAA
     751  CACGATTTTc GCGTCTTATC TGCAATCGGG CAGCGTTTCA TGGATGTATT
     801  ACGCCGACCG CATGATGGAG CTGCCAGCG GCGTGCTGGG GCGGCGACTC
     851  GGTACGATTT TGCTGCCGAC TTTGTCCAAA CACTCGGCAA ACCaAGATAC
     901  GGaACAGTTT TCCGCCCTGC TCGACTGGGG TTTGCGCCTG TGCATGCTgc
     951  TGACGCTGCC GCGGgcGGTC GGA CTGGCGG TGTGTGCTT cCCgCtGGTG
    1001  GCGACGCTGT TTATGTACCG CGwATTACG CTGTTTGACG CGCAGATGAC
    1051  GCAACACGCG CTGATTGCCT ATTCTTTCGG TTTAATCGGC TTAATCATGA
    1101  TTAAAGTGTT GGCACCGGCG TTCTATGCGC GGCAAAACAT CAAwAmGCCC
    1151  GTCAAATCG CCATCTTCAC GCTCATCTGC mCGCAGTTGA TGAACCTTGs
    1201  CTTTAYCGGC CCACTrAAC rCaSTCGGAC TTTTCGCTGC CATCGGTCTG
    1251  GGCGCGTGTA TCAATGCCGG ATTGTGTTT TACCTGTTGC GCAGACACGG
    1301  TATTTACCAA CTTGG.CAAG GGTGGGCAG CGTTCTT.AG CAAAAATGCT
    1351  GcTCTCGCTC GCCGTGA
  
```

This corresponds to the amino acid sequence <SEQ ID 114; ORF20>:

```

45      1  MNMLGALAKV GSLTMVSRVL GFVRDVIAR AFGAGMATDA FFVAFKLPNL
      51  LRRVFAEGAF AQAFVPILAE YKETRSKEAX EAFIRHVAGM LSFVLVIVTA
     101  LGILAAPWVI YVSAPSFAQD ADKFQLSIDL LRITFPYILL ISLSSFVGSV
     151  LNSYHKFGIP AFTPXFLNVS FIVEALFFVP YFDPVPTAXA WAVFVGILQ
     201  LXFQLPWLAK LGFLKLPKLS FKDAAVNRVM QMAPAILGV SVAQVSLVIN
     251  TIFASYLQSG SVSWMYADR MMELPSGVLG AALGTILLPT LSKHSANQDT
     301  EQFSALLDWG LRLCMLLTLP AAVGLAVLSF PLVATLFMYR XFTLFDAQMT
     351  QHALIAYSFG LIGLIMIKVL APGFYARQNI XXPVKIAIFT LICXQLMNLX
     401  FXGPLXXIGL SLAIGLGACI NAGLLFYLLR RHGIYQXPQG LGSVLXQKCC
     451  SRSP*
  
```

These sequences were elaborated, and the complete DNA sequence <SEQ ID 115> is:

```

55      1  ATGAATATGC TGGGAGCTTT GGCAAAAGTC GGCAGCCTGA CGATGGTGTG
      51  GCGCGTTTTG GGATTGTGTC GCGATACGGT CATTGCGCGG GCATTCGGCG
  
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101 CGGGTATGGC GACGGATGCG TTTTGTGCG CGTTCAAACCT GCCCAACCTG
 151 CTTCCGCCGCG TGTTTGCCTG GGGGCGGTTT GCCCAAGCGT TTGTGCCGAT
 201 TTGCGCCGAA TACAAGGAAA CGCGTTCAAA AGAGGCGGCG GAGGCTTTTA
 251 TCCGCCATGT GCGCGGGATG CTGTCGTTTG TACTGGTTAT CGTTACCGCG
 301 CTGGGCATAC TTGCCGCGCC TTGGGTGATT TATGTTTCCG CACCCGCTTT
 351 TGCCCAAGAT GCGGACAAAT TTCAGCTCTC CATCGATTG CTGCGGATTA
 401 CGTTTCCTTA TATATTATTG ATTTCCCTGT CTTCAATTGT CGGCTCGGTA
 451 CTCAATTCTT ATCATAAGTT CGGCATTCCG GCGTTTACGC CCACGTTTCT
 501 GAACGTGTCG TTTATCGTAT TCGCGCTGTT TTTCGTGCCG TATTTTCGATC
 551 CGCCCGTTAC CGCGCTGGCG TGGGCGGTCT TTGTCGGCGG CATTTTGCAA
 601 CTCGGCTTCC AACTGCCCTG GCTGGCGAAA CTGGGCTTTT TGAAACTGCC
 651 CAACATGAGT TTCAAAGATG CGGCGGTCAA CCGCGGTATG AAACAGATGG
 701 CGCCTGCGAT TTTGGGCGTG AGCGTGGCGC AGGTTTCTTT GGTGATCAAC
 751 ACGATTTTCG CGTCTTATCT GCAATCGGGC AGCGTTTCAT GGATGTATTA
 801 CGCCGACCGC ATGATGGAGC TGCCAGCGG CGTGCTGGGG GCGGCACTCG
 851 CACGATTTT GCTGCCGACT TTGTCCAAAC ACTCGGCAAA CCAAGATACG
 901 GAACAGTTTT CCGCCCTGCT CGACTGGGGT TTGCGCCTGT GCATGCTGCT
 951 GACGCTGCCG GCGGCGGTG GACTGGCGGT GTTGTCTGTC CCGCTGGTGG
 1001 CGACGCTGTT TATGTACCGC GAATTTACGC TGTTTGACGC GCAGATGACG
 1051 CAACACGCGC TGATTGCGTA TTCTTTCGGT TTAATCGGCT TAATCATGAT
 1101 TAAAGTGTG GCACCCGGCT TCTATGCGCG GCAAAACATC AAAACGCCCCG
 1151 TCAAAATCGC CATCTTCACG CTCATCTGCA CGCAGTTGAT GAACCTTGCC
 1201 TTTATCGGCC CACTGAAACA CGTCGGACTT TCGCTTGCCA TCGGCTCTGG
 1251 CGCGTGATC AATGCCGAT TGTGTTTTA CCTGTTGCGC AGACACGGTA
 1301 TTTACCAACC TGGCAAGGGT TGGGAGCGT TCTTAGCAA AATGCTGCTC
 1351 TCGCTCGCG TGATGTGCG GCGACTGTG GCGAGCGAGG CTACCTGCC
 1401 GTTTGAATGG GCGCACGCC GCGGAATCG GAAAGCGGG CAGCTCTGCA
 1451 TCTGATTCG CGTCGGCGC GGACTGTATT TCGCATCACT GCGGCTTTG
 1501 GGCTTCCGTC CGCGCCATTT CAAACGCGT GAAACTGA

30 This corresponds to the amino acid sequence <SEQ ID 116; ORF20-1>:

1 MNMLGALAKV GSILTMVSRVL GFVRDVIAR AFGAGMATDA FVFAFKLPNL
 51 LRRVFAEGAF AQAFVPILAE YKETRSKEAA EAFIRHVAGM LSFVLVIVTA
 101 LGILAAPWVI YVSAPGFAQD ADKFQLSIDL LRITFPYILL ISLSFVGSV
 151 LNSYHKFGIP AFTPTFLNVS FVFALEFFVP YFDPPTALA WAVFVGGLQ
 201 LGFLPWLAK LGFLKLPKLS FKDAAVNRVM KQMAPAILGV SVAQVSLVIN
 251 TIFASYLQSG SVSWMYADR MMELPSGVLG AALGTILLPT LSKHSANQDT
 301 EQFSALLDWG LRLCMLLTLP AAVGLAVLSF PLVATLFMYR EFTLFDAQMT
 351 QHALIAYSFG LIGLIMIKVL APGFYARQNI KTPVKIAIFT LICTQLMNL
 401 FIGPLKHVGL SLAIGLGACI NAGLLFYLLR RHGIYQPGKG WAAFLAKMLL
 451 SLAVMCGGLW AAQAYLPFEW AHAGGMRKAG QLCILIAVGG GLYFASLAAL
 501 GFRPRHFKEV EN*

Computer analysis of this amino acid sequence gave the following results:

Homology with the MviN virulence factor of *S. typhimurium* (accession number P37169)

ORF20 and MviN proteins show 63% aa identity in 440aa overlap:

45 Orf20 1 MNMLGALAKVGSILTMVSRVLGFVRDVIARAFGAGMATDAFFVFAFKLPNLLRRVFAEGAF 60
 MN+L +LA V S+TM SRVLGF RD ++AR FGAGMATDAFFVFAFKLPNLLRR+FAEGAF
 MviN 14 MNLLKSLAAVSSMTMFSRVLGFARDAIVARIFGAGMATDAFFVFAFKLPNLLRRIFAEGAF 73
 50 Orf20 61 AQAFVPILAIEYKETRSKEAXEAFIRHVAGMLSFVLVIVTALGILAAPWVIYVSAPGFAQD 120
 +QAFVPILAIEYK + +EA F+ +V+G+L+ L +VT G+LAAPWVI V+AP FA
 MviN 74 SQAFVPILAIEYKSKQGEATRIFVAYVSGLLTLALAVTVAGMLAAPWVIMVTAPGFADT 133
 55 Orf20 121 ADKFQLSIDLLRITFPYILLISLSFVGSVLSYHKFGIPFTXFLNVSFVFALEFFVP 180
 ADKF L+ LLRITFPYILLISL+S VG++LN+++F IPAF P FLN+S I FALF P
 MviN 134 ADKFALTTQLLRITFPYILLISLASLVGAILNTWNRFSIPAFPTFLNISMIGFALFAAP 193
 60 Orf20 181 YFDPPTAXAWAVFVGGLQLXFLPWLAKLGLKLPKLSFKDAAVNRVMKQMAPAILGV 240
 YF+PPV A AWAV VGG+LQL +QLP+L K+G L LP+++F+D RV+KQM PAILGV
 MviN 194 YFNPPVLALAWAVTVGGVLQVLYQLPYLKKIGMLVLPINFRDTGAMRVVKQMGPAAILGV 253
 Orf20 241 SVAQVSLVINTIFASYLQSGSVSWMYADRMELPSGVLGAALGTILLPTLSKHSANQDT 300
 SV+Q+SL+INTIFAS+L SGSVSWMYADR+ME PSGVLG ALGTILLP+LSK A+ +
 MviN 254 SVSQISLIINTIFASFLASGSVSWMYADRLMEFSPGVLGVALGTILLPSLSKSFASGNH 313

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Orf20 301 EQFSALLDWGLRLCMLLTLPAAVGLAVLSFPLVATLFMYRXFTLFDAQMTQHAIAYSFG 360
 +++ L+DWGLRLC LL LP+AV L +L+ PL +LF Y FT FDA MTQ ALIAYS G
 MviN 314 DEYCRIMDWGLRLCFLALPSAVALGILAKPLTVSLFQYGFATFADAMTQRALIAYSFG 373

5 Orf20 361 LIGLIMIKVLAPGFYARONIXXPVKIAIFTLICXQLMNLXFXXXXXXXXXXXXXXXXXXXCI 420
 LIGLI++KVLAPGFY+RQ+I PVKIAI TLI QLMNL F C+
 MviN 374 LIGLIVVKVLAPGFYSRQDIKTPVKIAIVTLIMTQLMNLAFIGPLKHAGLSLSIGLAACL 433

10 Orf20 421 NAGLLFYLLRRHGIYQXPQG 440
 NA LL++ LR+ I+ P G
 MviN 434 NASLLYWQLRKQNIPTQPG 453

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF20 shows 93.5% identity over a 447aa overlap with an ORF (ORF20a) from strain A of *N.*

15 *meningitidis*:

		10	20	30	40	50	60
orf20.pep		MNMLGALAKVGS	TMVSRVLGFVRD	TVIARAFGAGMAT	DAFFVAFKLPNLLRRVFAEGAF		
20 orf20a		MNMLGALVKVGS	TMVSRVLGFVRD	TVIARAFGAGMAT	DAFFVAFKLPNLLRRVFAEGAF		
		10	20	30	40	50	60
		70	80	90	100	110	120
orf20.pep		AQAFVPILA	EYKETSKEAXEAFIRHVAGMLSFVLVIVTALGILAAPWVIYVSAPSFQAD				
25 orf20a		AQAFVPILA	EYKETSKEATEAFIRHVAGMLSFVLVIVTALGILAAPWVIYVSAPGFAKD				
		70	80	90	100	110	120
		130	140	150	160	170	180
orf20.pep		ADKFQLSIDLLRITFPYILLISLS	SSFVGSVLNSYHKFGIPAF	TPXFLNVSFIVEALFFVP			
30 orf20a		ADKFQLSIDLLRITFPYILLISLS	SSFVGSVLNSYHKFSIPAF	TPFTFLNVSFIVEALFFVP			
		130	140	150	160	170	180
		190	200	210	220	230	240
35 orf20.pep		YFDPFPV	TAXAWAVFVG	GILQLXFPWLAKL	GFLKLPKLSFKDAAVNRVMQMAPAILGV		
orf20a		YFDPFPV	TALAWAVFVG	GILQLGFPWLAKL	GFLKLPKLSFKDAAVNRVMQMAPAILGV		
		190	200	210	220	230	240
		250	260	270	280	290	300
40 orf20.pep		SVAQVSLVINTIFASYLQSGSVSWMYADRM	MELPSGVLGAALGTILPTLSKHSANQDT				
orf20a		SVAQISLVINTIFASYLQSGSVSWMYADRM	MELPGGVLGAALGTILPTLSKHSANQDT				
		250	260	270	280	290	300
		310	320	330	340	350	360
orf20.pep		EQFSALLDWGLRLCMLLTLPAAVGLAVLSFPLVATLFMYRXFTLFDAQMTQHAIAYSFG					
50 orf20a		EQFSALLDWGLRXCMMLTLPAAVGMVAVLSFPLVATLFMYREFTLFDAQMTQHAIAYSFG					
		310	320	330	340	350	360
		370	380	390	400	410	420
orf20.pep		LIGLIMIKVLAPGFYARONIXXPVKIAIFTLICXQLMNLXFXG	PLXXIGLSLAIGLGACI				
55 orf20a		LIGLIMIKVLAPGFYARONIKTPVKIAIFTLICXQLMNLAFIGPLKHVGLSLAIGLGACI					
		370	380	390	400	410	420
		430	440	450			
orf20.pep		NAGLLFYLLRRHGIYQXPQGLG	SVLXQKCCSRSPX				
60 orf20a		NAGLLFYLLRRHGIYQPGKWA	AFLAKMLLSLAVMGGGLYAAQIWL	PFDDWAHAGGMQKAA			
		430	440	450	460	470	480

The complete length ORF20a nucleotide sequence <SEQ ID 117> is:

65 1 ATGAATATGC TGGGAGCTTT GCTAAAAGTC GGCAGCCTGA CGATGGTGTC
 51 GCGCGTTTGTG GGATTTGTGC GCGATACGGT CATTGCGCGC GCATTGCGCG
 101 CAGGCATGGC GACGGATGCG TTCTTTGTGC CGTTCAAACCTGCCCAACCTG

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151 CTTGCGCGCG TGTTCGCGGA GGGGCGGTTT GCCCAAGCGT TTGTGCGCAT
 201 TTTGGCGGAA TATAAGGAAA CGCGTCTTAA AGAGGCGACG GAGGCTTTTA
 251 TCCGCCATGT GGC GGCGGATG CTGTCGTTTG TACTGGTCAT CGTTACCGCG
 301 CTGGGCATAC TTGCGCGGCC TTGGGTGATT TATGTTCCG CACCCGGTTT
 351 TGCCAAAGAT GCCGACAAAT TTCAGCTCTC TATCGATTG CTGCGGATTA
 401 CGTTTCCTTA TATCTTATTG ATTTCACTTT CCTCTTTTGT CGGCTCGGTA
 451 CTCAATTCCT ATCATAAATT CAGCATTCCT GCGTTACGC CCACGTTCCCT
 501 GAACGTGTCG TTTATCGTAT TCGCGCTGTT TTTCTGCGG TATTTCGATC
 551 CTCCCCTTAC CGCGCTGGCT TGGGCGGTTT TTGTCGGCGG CATTTTGCAA
 601 CTCGGCTTCC AACTGCCCTG GCTGGCGAAA CTGGGTTTTT TGAAACTGCC
 651 CAAACTGAGT TTCAAAGATG CGGCGGTCAA CCGCGTGATG AAACAGATGG
 701 CGCCTGCGAT TTTGGGCGTG AGCGTGCGGC AGATTTCTTT GGTGATCAAC
 751 ACGATTTTCG CGTCTTATCT GCAATCGGGC AGCGTTTCAT GGATGTATTA
 801 CGCCGACCGC ATGATGGAAC TGCCCGGCGG CGTGCTGGGG GCGGCACTCG
 851 GTACGATTTT GCTGCCGACT TTGTCCAAAC ACTCGGCAAA CCAAGATACG
 901 GAACAGTTTT CCGCCCTGCT CCACTGGGGT TTGCGCNTGT GCATGCTGCT
 951 GACGCTGCCG GCGGCGGTCG GAATGGCGGT GTTGTCTGTC CCGCTGGTGG
 1001 CAACCTTGTT TATGTACCGA GAATTCACGC TGTTTGACGC GCAGATGACG
 1051 CAACACGCGC TGATTGCCTA TTCTTTCGGT TTAATCGGTT TAATCATGAT
 1101 TAAAGTGTTG GCGCCCGGCT TTTATGCGCG GCAAAACATC AAAACGCCCG
 1151 TCAAAATCGC CATCTTACG CTCATTTGCA CGCAGTTGAT GAACCTTGCC
 1201 TTTATCGGCC CACTGAAACA CGTCGGACTT TCGCTTGCCA TCGTCTGGG
 1251 CGCGTGATC AATGCCGGAT TGTTGTTTTA CCTGTTGCGC AGACACGGTA
 1301 TTTACCAACC TGGCAAGGGT TGGGCAGCGT TCTTGGCAAA AATGCTGCTC
 1351 TCGCTCGCCG TGATGGGAGG CGGCCTGTAT GCCGCCCCAA TCTGGCTGCC
 1401 GTTCGACTGG GCACACGCCG GCGGAATGCA AAAGCCGCCG CGGCTCTTCA
 1451 TCCTGATTGC CGTCGGCGGC GGACTGTATT TCGCATCACT GGCGGCTTTG
 1501 GGCTTCCGTC CGCGCCATTT CAAACGCGTG GAAAGCTGA

This encodes a protein having amino acid sequence <SEQ ID 118>:

30 1 MNMLGALVKV GSLTMVSRVL GFVRDVIAR AFGAGMATDA FFVAFKLPNL
 51 LRRVFAEGAF AQAFVPILAE YKETRSKEAT EAFIRHVAGM LSFVLVIVTA
 101 LGILAAPWVI YVSAPGFAKD ADKFQLSIDL LRITFPYILL ISLSSFVGSV
 151 LNSYHKFSIP AFTPTFLNVS FIVFALFFVP YFDPPTALAW WAVFVGILQ
 201 LGFQLPWLAK LGFLKLPKLS FKDAAVNRVM QMAPAILGV SVAQISLVIN
 35 251 TIFASYLQSG SVSWMYADR MMELPGGVLG AALGTILLPT LSKHSANQDT
 301 EQFSALLDWG LRXCMLLTLP AAVGMAVLSF PLVATLFMYR EFTLFDAQMT
 351 QHALIAYSFG LIGLIMIKVL APGFYARQNI KTPVKIAIFT LICTQLMNLA
 401 FIGPLKHVGL SLAIGLGACI NAGLLEYLLR RHGIYQPGKG WAAFLAKMLL
 451 SLAVMGGGLY AAQIWLFPDW AHAGGMQKAA RLFILIAVGG GLYFASLAAL
 40 501 GFRPRHFKRV ES*

ORF20a and ORF20-1 show 96.5% identity in 512 aa overlap:

		10	20	30	40	50	60
45	orf20a.pep	MNMLGALVKV	GSLTMVSRVL	GFVRDVIAR	AFGAGMATDA	FFVAFKLPNL	LRRVFAEGAF
	orf20-1	MNMLGALAKV	GSLTMVSRVL	GFVRDVIAR	AFGAGMATDA	FFVAFKLPNL	LRRVFAEGAF
		10	20	30	40	50	60
	orf20a.pep	AQAFVPILAE	YKETRSKEATE	EAFIRHVAGM	LSFVLVIVTA	LGILAAPWVI	YVSAPGFAKD
50	orf20-1	AQAFVPILAE	YKETRSKEAAE	EAFIRHVAGM	LSFVLVIVTA	LGILAAPWVI	YVSAPGFAQD
		70	80	90	100	110	120
55	orf20a.pep	ADKFQLSIDL	LRITFPYILL	ISLSSFVGSV	LNSYHKFSIP	AFTPTFLNVS	FIVFALFFVP
	orf20-1	ADKFQLSIDL	LRITFPYILL	ISLSSFVGSV	LNSYHKFGIP	AFTPTFLNVS	FIVFALFFVP
		130	140	150	160	170	180
60	orf20a.pep	YFDPPTALAW	AVFVGILQL	GFQLPWLAK	LGFLKLPKLS	FKDAAVNRVM	QMAPAILGV
	orf20-1	YFDPPTALAW	AVFVGILQL	GFQLPWLAK	LGFLKLPKLS	FKDAAVNRVM	QMAPAILGV
		190	200	210	220	230	240
65	orf20a.pep	SVAQISLVIN	TIFASYLQSG	SVSWMYADR	MMELPGGVLG	AALGTILLPT	LSKHSANQDT
		250	260	270	280	290	300

30 ORF20 shows 92.1% identity over a 454aa overlap with a predicted ORF (ORF20ng) from *N. gonorrhoeae*:

35	orf20.pep	MNMLGALAKVGSLTMVSRVLGFVRDVTIARAFAGMATDAFFVAFKLPNLLRRRVFAEGAF	60
	orf20ng	MNMLGALAKVGSLTMVSRVLGFVRDVTIARAFAGMATDAFFVAFKLPNLLRRRVFAEGAF	60
40	orf20.pep	AQAFVPILAEYKETRSKEAXEAFIRHVAGMLSFVLVIVTALGILAAPWVIYVSAPSFAQD	120
	orf20ng	AQAFVPILAEYKETRSKEATEAFIRHVAGMLSFVLIVVTALGILAAPWVIYVSAPGFTKD	120
45	orf20.pep	ADKFQLSIDLLRITFPYILLISLSSFVGSVLNSYHKFGIPAFTPXFLNVSFIVFALFFVP	180
	orf20ng	ADKFQLSISLLRITFPYILLISLSSFVGSILNSYHKFGIPAFTPTFLNISFIVFALFFVP	180
50	orf20.pep	YFDPPVTAXAWAVFVGILQLXFLQPLWLAKLGLKLPKLSFKDAAVNRVMKQMAPAILGV	240
	orf20ng	YFDPPVTALAWAVFVGILQLGFLQPLWLAKLGLKLPKLNFKDAAVNRVMKQMAPAILGV	240
55	orf20.pep	SVAQVSLVINTIFASYLQSGSVSWMYYADRMELPSGVLGAALGTILLPTLSKHSANQDT	300
	orf20ng	SVAQISLVINTIFASYLQSGSVSWMYYADRMELPGGVLGAALGTILLPTLSKHSANQDT	300
60	orf20.pep	EQFSALLDWGLRLCMLLTLPAAVGLAVLSFPLVATLFMYRXFTFLDQMTQHALIAYSFG	360
	orf20ng	EQFSALLDWGLRLCMLLTLPAAAGLAVLSFPLVATLFMYREFTFLDQMTQHALIAYSFG	360
65	orf20.pep	LIGLIMIKVLAPGFYARQNIXXPVKIAIFTLICXQLMNLXFXGPLXXIGLSLAIGLGACI	420
	orf20ng	LIGLIMIKVLASGFYARQNIKTPVKIAIFTLICQQLMNLAFIGPLKHAGLSLAIGLGACI	420
70	orf20.pep	NAGLLFYLLRRHGIYQFXQGLGSVLXQKCCSRSP	454
	orf20ng	NAGLLFFLFRKHGIYRPGQGLQPSWRKCCSRSP	454

An ORF20ng nucleotide sequence <SEQ ID 119> was predicted to encode a protein having amino acid sequence <SEQ ID 120>:

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1 MNMLGALAKV GSLTMVSRVL GFVRDVIAR AFGAGMATDA FFVAFKLPNL
 51 LRRVFAEGAF AQAFVPILAE YKETRSKEAT EAFIRHVAGM LSFVLIVVTA
 101 LGILAAPWVI YVSAPGFTKD ADKFQLSISL LRITFPYILL ISLSSFVSGSI
 151 LNSYHKFGIP AFTPTFLNIS FIVFALFFVP YFDPPTVATA WAVFVGGILQ
 201 LGFQLPWLAK LGFLKLPKLN FKDAAVNRVM KQMAPAILGV SVAQISLVIN
 251 TIFASYLQSG SVSWMYADR MMELPGGVLG AALGTILLPT LSKHSANQDT
 301 EQFSALLDWG LRLCMLLTLP AAAGLAVLSF PLVATLFMYR EFTLFDAQMT
 351 QHALIAYSFG LIGLIMIKVL ASGFYARQNI KTPVKIAIFT LICTQLMNL
 401 FIGPLKHAGL SLAIGLGACI NAGLLFFLLR KHGIYRPGQG LGQPSWRKCC
 451 SRSP*

Further DNA sequence analysis revealed the following DNA sequence <SEQ ID 121>:

1 ATGAATATGC TTGGAGCTTT GGCAAAAGTC GGCAGCCTGA CGATGGTGTG
 51 GCGCGTTTTG GGATTTGTGC GCGATACGGT CATTCGCGCG GCATTGCGCG
 101 CCGGTATGGC GACGGATGCG TTTTGTGTCG CGTTCAACT GCCCAACCTG
 151 CTTGCGCGCG TGTTCGCGA GGGGCGGTT GCCCAAGCGT TTGTGCGGAT
 201 TTTGGCGGAA TATAAGGAAA CGCGTCTTAA AGAGGCGA gAGGCTTTTA
 251 TCCGCCACGt tgcgggAatg CTGTCGTTG TGCTGATcgt cGttacCGCG
 301 CTGGGCATAC TTGCCGCGcc tTGGGTGATT TATGTTtccg CgcccGGCTT
 351 TACCAAGAC GCGGACAAAGT TCCAACCTTC CATCAGCGCT CTGCGGATTA
 401 CGTTTCCTTA TATATTATTG ATTTCTTTGT CTTCTTTTGT CGGCTCGATA
 451 CTAATTCCT ACCATAAGTT CGGCATTCCC GCGTTTACGC CCACGTTTTT
 501 AAACATCTCT TTTATCGTAT TCGCACTGTT TTTGCTGCCG TATTTGATC
 551 GCGCCGTTAC CGCGCTGGCG TGGGCGGTT TTGTCGCGCG TATTTTGCA
 601 CTCGGTTTCC AACTGCCGTG GCTGGCGAAA CTGGGCTTTT TGAAGTGGC
 651 CAACTGAAT TTCAAAGATG CGGCGGTCAA CCGCGTCATG AAACAGATGG
 701 CGCCTGCGAT TTTGGGCGTG agcgTGGCGC AAATTTCTTT GgttATCAAC
 751 ACGATTTTCG CGTCTTATCT GCAATCGGCG AGCGTTTCAT GGATGAtta
 801 cgCCGACCGC ATGATGGAGc tgcgcccGGG CGTGTGGGG GCTGCACTCG
 851 GTACAATTTT GCTGCCGACT TTGTCCAAAC ACTCGGCAAA CCAAGATACG
 901 GAACAGTTTT CCGCCCTGCT CGACTGGGGT TTGCGCCTGT GCATGCTGT
 951 GACGCTGCCG GCGGCGGccg GACTGGCGGT ATGTGCTTC CCGCTGGTGG
 1001 CGACGCTGTT TATGTACCGA GAATTCACGC TGTTCGACGC ACAATGACG
 1051 CAACACGCGC TGATTGCCTA TTCTTTCGGT TTAATCGGTT TAATTATGAT
 1101 TAAAGTGTG GCATCCGGCT TTTATGCGCG GCAAACATC AAAACGCCG
 1151 TCAAATCGC CATCTTCACG CTCATCTGCA CGCAGTTGAT GAACCTCGCC
 1201 TTTATCGGTC CGTTGAAACA CGCGGGCTT TCGCTCGCCA TCGCCTGGG
 1251 CCGGTGCATC AACGCCGGAT TGTGTTCTT CCGTTGCGC AAACACGGTA
 1301 TTTACCGGCC cggcaggggt tggcgcggt TCTTGGCGAA AATGCTGCTC
 1351 GCGCTCGCCG TGATGTGCGG CGGACTGTGG GCGGCGCAGG CTTGCTGCC
 1401 GTTCGAATGG GCGCACGCCG GCGGAATGCG GAAAGCGGG CAGCTCTGCA
 1451 TCCTGATTGC CGTCGGCGGC GGAATGTATT TCGCATCTCT GCGCGCTTTG
 1501 GGCTTCCGTC CGCGCCATTT CAAACGCGTG GAAAGCTGA

This encodes the following amino acid sequence <SEQ ID 122; ORF20ng-1>:

1 MNMLGALAKV GSLTMVSRVL GFVRDVIAR AFGAGMATDA FFVAFKLPNL
 51 LRRVFAEGAF AQAFVPILAE YKETRSKEAT EAFIRHVAGM LSFVLIVVTA
 101 LGILAAPWVI YVSAPGFTKD ADKFQLSISL LRITFPYILL ISLSSFVSGSI
 151 LNSYHKFGIP AFTPTFLNIS FIVFALFFVP YFDPPTVATA WAVFVGGILQ
 201 LGFQLPWLAK LGFLKLPKLN FKDAAVNRVM KQMAPAILGV SVAQISLVIN
 251 TIFASYLQSG SVSWMYADR MMELRRGVLG AALGTILLPT LSKHSANQDT
 301 EQFSALLDWG LRLCMLLTLP AAAGLAVLSF PLVATLFMYR EFTLFDAQMT
 351 QHALIAYSFG LIGLIMIKVL ASGFYARQNI KTPVKIAIFT LICTQLMNL
 401 FIGPLKHAGL SLAIGLGACI NAGLLFFLLR KHGIYRPGRG WAAFLAKMLL
 451 ALAVMCGGLW AAQACLPEW AHAGGMRKAG QLCILIAVGG GLYFASLAAL
 501 GFRPRHFKRV ES*

55 ORF20ng-1 and ORF20-1 show 95.7% identity in 512 aa overlap:

	10	20	30	40	50	60
orf20-1.pep	MNMLGALAKV	GSLTMVSRVL	GFVRDVIAR	AFGAGMATDA	FFVAFKLPNL	LRRVFAEGAF
orf20ng-1	MNMLGALAKV	GSLTMVSRVL	GFVRDVIAR	AFGAGMATDA	FFVAFKLPNL	LRRVFAEGAF
60	10	20	30	40	50	60
	70	80	90	100	110	120
orf20-1.pep	AQAFVPILAE	YKETRSKEA	EAFIRHVAG	MLSFVLIV	TALGILAAP	WVIYVSAP
orf20ng-1	AQAFVPILAE	YKETRSKEA	EAFIRHVAG	MLSFVLIV	TALGILAAP	WVIYVSAP
65						

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		70	80	90	100	110	120
		130	140	150	160	170	180
5	orf20-1.pep	ADKFQLSIDLLRITFPYILLISLSSSVGSLNSYHKEGIPAFPTPTFLNVSFIVFALFFVP					
	orf20ng-1	ADKFQLSISLLRITFPYILLISLSSSVGSLNSYHKEGIPAFPTPTFLNISFIVFALFFVP					
		130	140	150	160	170	180
		190	200	210	220	230	240
10	orf20-1.pep	YFDPVPTALAWAVFVGILQLGFQLPWLAKLGLKLPKLSFKDAAVNRVMKQMAPAILGV					
	orf20ng-1	YFDPVPTALAWAVFVGILQLGFQLPWLAKLGLKLPKLNFKDAAVNRVMKQMAPAILGV					
		190	200	210	220	230	240
		250	260	270	280	290	300
15	orf20-1.pep	SVAQVSLVINTIFASYLQSGSVSWMYADRMMLPSGVLGAALGTILLPTLSKHSANQDT					
	orf20ng-1	SVAQISLVINTIFASYLQSGSVSWMYADRMMLRRGVLGAALGTILLPTLSKHSANQDT					
		250	260	270	280	290	300
		310	320	330	340	350	360
20	orf20-1.pep	EQFSALLDWGLRLCMLLTLPAAVGLAVLSFPLVATLFMYREFTLFDAQMTQHAIAYSFG					
	orf20ng-1	EQFSALLDWGLRLCMLLTLPAAGLAVLSFPLVATLFMYREFTLFDAQMTQHAIAYSFG					
25		310	320	330	340	350	360
		370	380	390	400	410	420
30	orf20-1.pep	LIGLIMIKVLAPGFYARQNIKTPVKIAIFTLICTQLMNLAFIGPLKHVGLSLAIGLGACI					
	orf20ng-1	LIGLIMIKVLASGFYARQNIKTPVKIAIFTLICTQLMNLAFIGPLKHAGLSLAIGLGACI					
		370	380	390	400	410	420
		430	440	450	460	470	480
35	orf20-1.pep	NAGLLFYLLRRHGIYPGKGWAAFLAKMLLSLAVMCGGLWAAQAYLPFEWAHAGGMRKAG					
	orf20ng-1	NAGLLFFLLRKHGIYRPGRWAAFLAKMLLALAVMCGGLWAAQACLPFEWAHAGGMRKAG					
		430	440	450	460	470	480
		490	500	510			
40	orf20-1.pep	QLCILIAVGGGLYFASLAALGFRPRHFKRVENX					
	orf20ng-1	QLCILIAVGGGLYFASLAALGFRPRHFKRVESX					
		490	500	510			

In addition, ORF20ng-1 shows significant homology with a virulence factor of *S.typhimurium*:

45	sp P37169 MVIN_SALTY_VIRULENCE_FACTOR_MVIN_pir S40271 mviN protein - Salmonella typhimurium gi 438252 (Z26133) mviB gene product [Salmonella typhimurium] gnl PID d1005521 (D25292) ORF2 [Salmonella typhimurium] Length = 524 Score = 1573 (750.1 bits), Expect = 1.1e-220, Sum P(2) = 1.1e-220 Identities = 309/467 (66%), Positives = 368/467 (78%)
50	Query: 1 MNMLGALAKVGSMTMVSRLVGFVRDVIARAFAAGMATDAFFVAFKLPNLLRRVFAEGAF 60 MN+L +LA V S+TM SRVLGF RD ++AR FGAGMATDAFFVAFKLPNLLRR+FAEGAF Sbjct: 14 MNLLKSLAAVSSMTMFSRVLGFARDAIVARIFGAGMATDAFFVAFKLPNLLRRIFAEGAF 73
55	Query: 61 AQAQFVPILAIEYKETSKEATEAFIRHVAGMLSFVLIVVTALGILAAPWVIYVSAPGETKD 120 +QAQFVPILAIEYK + +EAT F+ +V+G+L+ L VVT G+LAAPWVI V+APGF Sbjct: 74 SQAQFVPILAIEYKSKQGEETRIFVAYVSGLLTLALAVVTVAGMLAAPWVIMVTAPGFADT 133
60	Query: 121 ADKFQLSISLLRITFPYILLISLSSSVGSLNSYHKEGIPAFPTPTFLNISFIVFALFFVP 180 ADKF L+ LLRITFPYILLISL+S VG+ILN++++F IPAF PTFNLIS I FALF P Sbjct: 134 ADKFALTQQLRITFPYILLISLASLVGAILNTWNRFSIPAFAPTFLNISMIGFALFAAP 193
65	Query: 181 YFDPVPTALAWAVFVGILQLGFQLPWLAKLGLKLPKLNFKDAAVNRVMKQMAPAILGV 240 YF+PPV ALAWAV VGG+LQL +QLP+L K+G L LP++NF+D RV+KQM PAAILGV Sbjct: 194 YFNFPVLALAWAVTVGGVLQLVYQLPYLKKIGMLVLPINFRDGTGAMRVVKQMGPAAILGV 253
70	Query: 241 SVAQISLVINTIFASYLQSGSVSWMYADRMMLRRGVLGAALGTILLPTLSKHSANQDT 300 SV+QISL+INTIFAS+L SGSVSWMYADR+ME GVLG ALGTILLP+LSK A+ + Sbjct: 254 SVSQISLIINTIFASFLASGSVSWMYADRLMEFPGVLGVALGTILLP+LSKSFASGNH 313

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Query: 301 EQFSALLDWGLRLCMLLTLPAAAGLAVLSFPLVATLFMYREFTLFDAQMTQHAIAYSFG 360
 +++ L+DWGLRLC LL LP+A L +L+ PL +LF Y +FT FDA MTQ ALIAYS G
 Sbjct: 314 DEYCRLLMDWGLRLCFLALPSAVALGILAKPLTVSLFQYGKFTAFDAAMTORALIAYSFG 373

5 Query: 361 LIGLIMIKVLASGFYARQNIKTPVKIAIFTLICTQLMNLAFIGPLKHAGLSLAIGLGACI 420
 LIGLI++KVLG GFY+RQ+IKTPVKIAI TLI TQLMNLAFIGPLKHAGLSL+IGL AC+
 Sbjct: 374 LIGLIVVKVLAPGFYSRQDIKTPVKIAIVTLIMTQLMNLAFIGPLKHAGLSLSIGLAACL 433

10 Query: 421 NAGLLFFLLRKHGIYRPGRGWXXXXXXXXXXXXXVMCGGLWAAQACLP 467
 NA LL++ LRK I+ P GW VM L+ +P
 Sbjct: 434 NASLLYWQLRKQNIPTPQPGWMWFLMRLIISVLVMAAVLFGVLHIMP 480

Score = 70 (33.4 bits), Expect = 1.1e-220, Sum P(2) = 1.1e-220
 Identities = 14/41 (34%), Positives = 23/41 (56%)

15 Query: 469 EWAHAGGMRKAGQLCILIAVGGGLYFASLAALGFRPRHFKR 509
 EW+ + + +L ++ G YFA+LA LGF+ + F R
 Sbjct: 481 EWSQGSMLWRLRLMAVVIAGIAAYFAALAVLGFKVKEFVR 521

- 20 Based on this analysis, including the homology with a virulence factor from *S.typhimurium*, it is predicted that these proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 15

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 123>:

25 1 atGATTAAAA TCAAAAAAGG TCTAAACCTG CCCATCGCGG GCAGACCGGA
 51 GCAAGCCGTT tACGACGGCC CGGCCaTTAC CGAAGtCGCG TTGCTTGCGG
 101 AAGAATATGC CGGTATGCGC CCCTCGATGA AAGTCAAGGA AGGCGATGCC
 151 GTCAAAAAAG GCCAAGTGCT GTTTGAAGAC AAAAAGAATC CGGGCGTGCT
 201 GTTACTGCG CCGGCTTCAG GcAAAATCGC CGCGATTAC CGTGGCGAAA
 30 AGCGCGTACT TCAGTCAGTC GTGATTGCCG TTGAAGGCAA CGACGAAATC
 301 GAGTTTGAAC GCTACGCACC TGAAGCGCTG GCAAACCTAA GCGGCGAAGA
 351 AGTGGCGCCG AACCTGATCC AATCCGGTTT GTGGACTGCG CTGCGCACCC
 401 GTCCGTTTCA CAAAATTCCT GCCGTCGATG CCGAGCCGTT CGCCATCTTC
 451 GTCAATGCCA tGGACACCAA TCCG..

- 35 This corresponds to the amino acid sequence <SEQ ID 124; ORF22>:

1 MIKIKKGLNL PIAGRPEQAV YDGPATEVA LLGEEYAGMR PSMKVKEGDA
 51 VKKGQVLFED KKNPGVVFTA PASGKIAAIH RGEKRVLQSV VIAVEXNDEI
 101 EFERYAPEAL ANLSGEEVRR NLIQSGLWTA LRTRPFSKIP AVDAEPFAIF
 151 VNAMDTPN..

- 40 Further work revealed the complete nucleotide sequence <SEQ ID 125>:

1 ATGATTAAAA TCAAAAAAGG TCTAAACCTG CCCATCGCGG GCAGACCGGA
 51 GCAAGCCGTT TACGACGGCC CGGCCATTAC CGAAGTCGCG TTGCTTGCGG
 101 AAGAATATGC CGGTATGCGC CCCTCGATGA AAGTCAAGGA AGGCGATGCC
 151 GTCAAAAAAG GCCAAGTGCT GTTTGAAGAC AAAAAGAATC CGGGCGTGCT
 45 201 GTTACTGCG CCGGCTTCAG GCAAAATCGC CGCGATTAC CGTGGCGAAA
 251 AGCGCGTACT TCAGTCAGTC GTGATTGCCG TTGAAGGCAA CGACGAAATC
 301 GAGTTTGAAC GCTACGCACC TGAAGCGCTG GCAAACCTAA GCGGCGAAGA
 351 AGTGGCGCCG AACCTGATCC AATCCGGTTT GTGGACTGCG CTGCGCACCC
 401 GTCCGTTTCA CAAAATTCCT GCCGTCGATG CCGAGCCGTT CGCCATCTTC
 50 451 GTCAATGCCA TGGACACCAA TCCGCTGGCT GCCGACCCTA CGGTCATTAT
 501 CAAAGAAGCC GCCGAGGATT TCAAACGCGG CCTGTTGGTA TTGAGCCGTT
 551 TGACCGAAGC CAAAATCCAT GTTTGTAAGG CAGCTGGCGC AGACGTGCCG
 601 TCTGAAATG CTGCCAACAT CGAAACACAT GAATTCGGCG GCCCGCATCC
 651 TGCCGGTTTG AGTGGCACGC ACATTCATT TATCGAGCCG CCGGCGCGGA
 701 ATAAACCGT GTGGACCATC AATTATCAAG ATGTAATTAC CATTGGCCGT
 751 TTGTTTGCAA CAGGCCGCT GAACACCGAG CGCGTGATTG CCCTAGGTGG
 801 TTCTCAAGTC AACAAACCGC GCCTCTTGG TACCGTTTTG GGTGCGAAG
 851 TATCGCAAAT TACTGCGGGC GAATTGGTTG ACACAGACAA CCGCGTGATT
 901 TCCGGTTCCG TATTGAACGG CGCGATTACA CAAGGCGCGC ACGATTATTT

5 951 GGGACGCTAC CACAATCAGA TTTCCGTTAT CGAAGAAGGC CGCAGCAAAG
1001 AGCTGTTTCGG CTGGGTTGCG CCGCAGCCGG ACAAATACTC CATCACGCGT
1051 ACAACCCTCG GCCATTTTCCT GAAAAACAAA CTCTTCAAGT TCAACACAGC
1101 CGTCAACGGC GGCACCCGCG CCATGGTGCC GATTGGTACT TACGAGCGCG
1151 TGATGCCCTT GGATATCCTG CCCACCCTGC TTTTGCGCGA TTTAATCGTC
1201 GGCATACCG ACAGCGCGCA GGCATTGGGT TGCTTGAAT TGGACGAAGA
1251 AGACCTCGCT TTGTGCAGCT TCGTCTGCC GGGCAAATAC GAATACGGCC
1301 CGCTGTTGCG CAAAGTGCTG GAAACCATTG AGAAGGAAGG CTGA

This corresponds to the amino acid sequence <SEQ ID 126; ORF22-1>:

10 1 MIKIKKGLNL PIAGRPEQAV YDGPATEVA LLGEEYAGMR PSMKVKEGDA
51 VKKQVLFED KKNPGVVFTA PASGKIAAIH RGEKRVLSQSV VIAVEGNDEI
101 EFERYAPEAL ANLSGEEVRR NLIQSGLWTA LRTRPFSKIP AVDAEPFAIF
151 VNAMDTNPLA ADPTVIIKEA AEDFKRGLLV LSRLTERKIH VCKAAGADV
201 SENAANIETH EFGGPHFAGL SGTHHFIEP VGANKTVWTI NYQDVITIGR
15 251 LFATGRLNTE RVIALGGSQV NKPRLLRTVL GAKVSQITAG ELVDTDNRVI
301 SGSVLNGAIT QGAHDYLGRI HNQISVIEEG RSKELFGWVA PQPDKYSITR
351 TTLGHFLKNK LFKFNTAVNG GDRAMVPIGT YERVMPLDIL PTLRLRLDILV
401 GDTDSAQALG CLELDEEDLA LCSFVCPGKY EYGPLLRKVL ETIEKEG*

Further work identified the corresponding gene in strain A of *N.meningitidis* <SEQ ID 127>:

20 1 ATGATTAAAA TCAAAAAGG TCTAAACCTG CCCATCGCGG GCAGACCGGA
51 GCAAGTCATT TATGACGGGC CCGTCATTAC CGAAGTCGCG TTGCTTGGCG
101 AAGAATATGC CCGTATGCGC CCCTNGATGA AAGTCAAGGA AGGCGATGCC
151 GTCAAAAAG GCCAAGTGCT GTTGAAGAC AAAAAGNATC CGGGCGTGGT
201 GTTTACCGCG CCNGTTTCAG GCAAAATCGC CGCCATCCAT CGCGGCGAAA
25 251 AGCGCGTACT TCAGTCGGTC GTGATTGCCG TTGAAGGCAA CGACGAAATC
301 GAGTTCGAAC GCTACGCGCC CGAAGCGTTG GCAAACTTAA GCGGCGANGA
351 ANTNNNGNGC AATCTGATCC AATCCGGTTT GTGGAAGTGC CTGCGTANCC
401 GTCCGTTTCA CAAAATCCCT GCCGTCGATG CCGAGCCGTT CGCCATCTTC
451 GTCGAATGCGA TGGACACCAA TCCGCTNGCG GCAGACCTG TGGTGTGAT
30 501 CAAAGAAGCC GNCGANGATT TCAGACGANG TNTGCTGGTA TTGAGCCGTT
551 TGACCGAGCG TAAATCCAT GTGTGTAAGG CAGCTGGCGC AGACGTGCCG
601 TCTGAAAATG CTGCCAACAT CGAAACACAT GAATTCGGCG GCCCGCATCC
651 GCGCGGTTTG AGTGGCACGC ACATTCATTT CATTGAGCCG GTCCGTGCAA
701 ACAAACCGT TTGGACCATC AATTATCAAG ATGTAATTGC CATCGGACGT
35 751 TTGTTTGCAA CAGGCCGTCT GAACACCGAG CGCGTGATG CTTTGGGTGG
801 TTCTCAAGTC AACAAACAC GCCTCTTGCG TACCGTTTGG GGTGCGAAAG
851 TATCGCAAAT TACTGCGGGC GAATTGGTTG ACGCAGACAA CCGCGTGATT
901 TCCGGTTTCG TATTGAACGG CGCGATTACA CAAGGCGCGC ACGATTATTT
951 GGGACGCTAC CACAATCAGA TTTCCGTTAT CGAAGAAGGC CGCAGCAAAG
40 1001 AGCTGTTTCGG CTGGGTTGCG CCGCAGCCGG ACAAATACTC CATCACGCGT
1051 ACGACCCTCG GCCATTTTCCT GAAAAACAAA CTCTTCAAGT TCACGACAGC
1101 CGTCAACGGT GGCACCCGCG CCATGGTGCC GATTGGTACT TACGAGCGCG
1151 TAATGCCGCT AGACATCCTG CCTACCCTGC TTTTGCGCGA TTTAATCGTC
1201 GGCATACCG ACAGCGCGCA AGCATTGGGT TGCTTGAAT TGGACGAAGA
45 1251 AGACCTCGCT TTGTGCAGCT TCGTCTGCC GGGCAAATAC GAATANGGCC
1301 CGCTGTTGCG TAAGTGCTG GAAACCNTTG AGAAGGAAGG CTGA

This encodes a protein having amino acid sequence <SEQ ID 128; ORF22a>:

50 1 MIKIKKGLNL PIAGRPEQVI YDGPVITEVA LLGEEYAGMR PXMKVKEGDA
51 VKKQVLFED KKNPGVVFTA PVSCKIAAIH RGEKRVLSQSV VIAVEGNDEI
101 EFERYAPEAL ANLSGXEXXX NLIQSGLWTA LRTRPFSKIP AVDAEPFAIF
151 VNAMDTNPLA ADPVVVIKEA XXDFRXXLV LSRLTERKIH VCKAAGADV
201 SENAANIETH EFGGPHFAGL SGTHHFIEP VGANKTVWTI NYQDVIAIGR
251 LFATGRLNTE RVIALGGSQV NKPRLLRTVL GAKVSQITAG ELVDADNRVI
301 SGSVLNGAIT QGAHDYLGRI HNQISVIEEG RSKELFGWVA PQPDKYSITR
55 351 TTLGHFLKNK LFKFTTAVNG GDRAMVPIGT YERVMPLDIL PTLRLRLDILV
401 GDTDSAQALG CLELDEEDLA LCSFVCPGKY EXGPLLRKVL ETXEKEG*

The originally-identified partial strain B sequence (ORF22) shows 94.2% identity over a 158aa overlap with ORF22a:

60 orf22.pep 10 20 30 40 50 60
MIKIKKGLNLPIAGRPEQAVYDGPATEVALLGEEYAGMRPSMKVKEGDAVKKQVLFED
orf22a MIKIKKGLNLPIAGRPEQVIYDGPVITEVALLGEEYAGMRPXMKVKEGDAVKKQVLFED

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		10	20	30	40	50	60
5	orf22.pep	70	80	90	100	110	120
		KKNPGVVFTAPASGKIAAIHRGEKRVLSVVI	AVEXNDEIEFERYAPEALANLSGEEVRR				
	orf22a	70	80	90	100	110	120
		KKXPGVVFTAPVSGKIAAIHRGEKRVLSVVI	AVEGNDEIEFERYAPEALANLSGXEXXX				
10	orf22.pep	130	140	150			
		NLIQSGLWTALRTRPFSKIPAVDAEPFAIFV	NAMDTNP				
	orf22a	130	140	150	160	170	180
		NLIQSGLWTALRTRPFSKIPAVDAEPFAIFV	NAMDTNP	LAADPVVVIKEAXXD	FRRXXLV		

The complete strain B sequence (ORF22-1) and ORF22a show 94.9% identity in 447 aa overlap:

15	orf22a.pep	10	20	30	40	50	60
		MIKIKKGLNLPIAGRPEQVIYDGPVITEV	ALLGEEYAGMRPXMVKVKEGDAVKKGQVLFED				
	orf22-1	10	20	30	40	50	60
		MIKIKKGLNLPIAGRPEQAVYDGPATTEV	ALLGEEYAGMRPSMKVKEGDAVKKGQVLFED				
20	orf22a.pep	70	80	90	100	110	120
		KKXPGVVFTAPVSGKIAAIHRGEKRVLSVVI	AVEGNDEIEFERYAPEALANLSGXEXXX				
25	orf22-1	70	80	90	100	110	120
		KKNPGVVFTAPASGKIAAIHRGEKRVLSVVI	AVEGNDEIEFERYAPEALANLSGEEVRR				
30	orf22a.pep	130	140	150	160	170	180
		NLIQSGLWTALRTRPFSKIPAVDAEPFAIFV	NAMDTNP	LAADPVVVIKEAXXD	FRRXXLV		
	orf22-1	130	140	150	160	170	180
		NLIQSGLWTALRTRPFSKIPAVDAEPFAIFV	NAMDTNP	LAADPTVIKEAAED	FKRGLLV		
35	orf22a.pep	190	200	210	220	230	240
		LSRLTERKIHVCKAAGADVPSENAANIETHE	FGGPHPAGLSGTHIHFI	EPVGANKT	VWTI		
	orf22-1	190	200	210	220	230	240
		LSRLTERKIHVCKAAGADVPSENAANIETHE	FGGPHPAGLSGTHIHFI	EPVGANKT	VWTI		
40	orf22a.pep	250	260	270	280	290	300
		NYQDVIAIGRLFATGRLNTERVIALGGSQV	NKPRLRTVLGAKVSQITAGELVDADNRVI				
	orf22-1	250	260	270	280	290	300
		NYQDVITIGRLFATGRLNTERVIALGGSQV	NKPRLRTVLGAKVSQITAGELVD	TADNRVI			
45	orf22a.pep	310	320	330	340	350	360
		SGSVLNGAITQGAHDYLGRYHNQISVIEEGR	SKELFGWVAPQPKYSITRTTLGHFLKNK				
	orf22-1	310	320	330	340	350	360
		SGSVLNGAITQGAHDYLGRYHNQISVIEEGR	SKELFGWVAPQPKYSITRTTLGHFLKNK				
50	orf22a.pep	370	380	390	400	410	420
		LFKETTAVNGGDRAMVPIGTYERVMPLDILP	TLLRLDLIVGDTDSAQALGCLELDEEDLA				
55	orf22-1	370	380	390	400	410	420
		LFKENTAVNGGDRAMVPIGTYERVMPLDILP	TLLRLDLIVGDTDSAQALGCLELDEEDLA				
60	orf22a.pep	430	440				
		LCSFVCPGKYEXGPLL	RKVLETXEKEGX				
	orf22-1	430	440				
		LCSFVCPGKYEYGPLL	RKVLETIEKEGX				

Further work identified a partial gene sequence <SEQ ID 129> from *N.gonorrhoeae*, which encodes the following amino acid sequence <SEQ ID 130; ORF22ng>:

65	1	MIKIKKGLNL	PIAGRPEQVI	YDGPATEVA	LLGEEYVGM	PSMKIKEGEA
	51	VKKGQVLFED	KKNPGVVFTA	PASGKIAAIH	RGEKRVLSV	VIAVEGNDEI
	101	EFERYVPEAL	AKLSSEKVR	R NLIQSGLWTA	LRTRPFSKIP	AVDAEPFAIF

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45 overlap with ORF22ng:

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orf22-1.pep 10 20 30 40 50 60
MIKIKKGLNLPIAGRPEQAVYDGPATTEVALLGEEYAGMRPSMKVKEGDAVKKGOVLFEED

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	orf22ng-1	: : : :	
		MIKIKKGLNLPIAGRPEQVIYDGPATEVALLGEEYVGM RPSMKIKEGEAVKKGQVLFED	
		10 20 30 40 50 60	
5	orf22-1.pep	70 80 90 100 110 120	
		KKNPGVVFTAPASGKIAAIHRGEKRVLSVVI AVEGNDEIEFERYAPEALANLSGEEVRR	
	orf22ng-1	: : : :	
		KKNPGVVFTAPASGKIAAIHRGEKRVLSVVI AVEGNDEIEFERYVPEALAKLSSEKVR	
		70 80 90 100 110 120	
10	orf22-1.pep	130 140 150 160 170 180	
		NLIQSGLWTALRTRPFSKIPAVDAEPFAIFVNAMDTNPLAADPTVIIKEAAEDFKRGLLV	
	orf22ng-1	: : : :	
		NLIQSGLWTALRTRPFSKIPAVDAEPFAIFVNAMDTNPLAADPTVIIKEAAEDFKRGLLV	
		130 140 150 160 170 180	
15	orf22-1.pep	190 200 210 220 230 240	
		LSRLTERKIHVCKAAGADVPSENAANIETHEFGGPHPAGLSGTHIFIEPVGANKTVWTI	
	orf22ng-1	: : : :	
		LSRLTERKIHVCKAAGADVPSENAANIETHEFGGPHPAGLSGTHIFIEPVGANKTVWTI	
		190 200 210 220 230 240	
20	orf22-1.pep	250 260 270 280 290 300	
		NYQDVITIGRLFATGRLNTERVIALGGSQV NKPRLRLTVLGAKVSQITAGELVDNDRVI	
	orf22ng-1	: : : :	
		NYQDVIAIGRLFVTGRLNTERVVALGGLQV NKPRLRLTVLGAKVSQITAGELVDADNRVI	
		250 260 270 280 290 300	
25	orf22-1.pep	310 320 330 340 350 360	
		SGSVLNGAITQGAHDYLG RYHNQISVIEEGRSKELFGWVAPQDPKYSITRTTLGHFLKKNK	
	orf22ng-1	: : : :	
		SGSVLNGAIAQGAHDYLG RYHNQISVIEEGRSKELFGWVAPQDPKYSITRTTLGHFLKKNK	
		310 320 330 340 350 360	
30	orf22-1.pep	370 380 390 400 410 420	
		LFKFNTAVNGGDRAMVPIGT YERVMPLDILPTLLLRDLIVGDTDSAQALGCLELDEEDLA	
	orf22ng-1	: : : :	
		LFKFNTAVNGGDRAMVPIGT YERVMPLDILPTLLLRDLIVGDTDSAQALGCLELDEEDLA	
		370 380 390 400 410 420	
35	orf22-1.pep	430 440	
		LCSFVCPGKYEGP LLRKVLETIEKEGX	
	orf22ng-1	: : : :	
		LCSFVCPGKYEGP LLRKVLETIEKEGX	
		430 440	
40			
45			

Computer analysis of these sequences gave the following results:

Homology with 48kDa outer membrane protein of *Actinobacillus pleuropneumoniae* (accession number U24492).

ORF22 and this 48kDa protein show 72% aa identity in 158aa overlap:

50	Orf22	1	MIKIKKGLNLPIAGRPEQAVYDGPATEVALLGEEYAGMRPSMKVKEGDAVKKGQVLFED	60
			MI IKKGL+LPIAG P Q +++G + EVA+LGEEY GMRPSMKV+EGD VKKGQVLFED	
	48kDa	1	MITIKKGLDLPIAGTPAQVIHNGNTVNEVAMLGEEYVGM RPSMKVREGDVVKKGQVLFED	60
	orf22	61	KKNPGVVFTAPASGKIAAIHRGEKRVLSVVI AVEGNDEIEFERYAPEALANLSGEEVRR	120
			KKNPGVVFTAPASG + I+RGEKRVLSVVI VE +++I F RY LA+LS E+V++	
55	48kDa	61	KKNPGVVFTAPASGTVVTTINRGEKRVLSVVIKVEGDEQITFTRYEAAQLASLSAEQVKQ	120
	orf22	121	NLIQSGLWTALRTRPFSKIPAVDAEPFAIFVNAMDTNP	158
			NLI+SGLWTA RTRPFSK+PA+DA P +IFVNAMDTNP	
60	48kDa	121	NLIESGLWTAFRTRPFSKVPALDAIPSSIFVNAMDTNP	158

ORF22a also shows homology to the 48kDa *Actinobacillus pleuropneumoniae* protein:

gi|1185395 (U24492) 48 kDa outer membrane protein [Actinobacillus pleuropneumoniae]
Length = 449

65 Score = 530 bits (1351), Expect = e-150

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Identities = 274/450 (60%), Positives = 323/450 (70%), Gaps = 4/450 (0%)

5 Query: 1 MIKIKKGLNLPIAGRPEQVIYDGPVITEVALLGEEYAGMRPXMVKVEGDAVKKGQVLFED 60
 Sbjct: 1 MITIKKGLDLPIAGTPAQVIHNGNTVNEVAMLGEEYVGMPSMKVREGDVVKKGQVLFED 60

10 Query: 61 KKKPGVVFTAPVSGKIAAIHRGEKRVLSVVI AVEGNDEIEFERYAPEALANLSGXEXXX 120
 Sbjct: 61 KKNPGVVFTAPASGTVVTINRGEKRVLSVVIKVEGDEQITFTRYEAAQLASLSAEQVKQ 120

15 Query: 121 NLIQSGLWLTALRXRPFSKIPAVDAEPFAIFVNAMDTNPLAADPVVVIKEAXXDFRXXLV 180
 Sbjct: 121 NLIESGLWTAFRTRPFSKVPALDAIPSSIFVNAMDTNPLAADPEVVLKEYETDFKDGLTV 180

20 Query: 238 WTINYQDVIAIGRLFATGRNLNTERVIALGGSQVKNPRLRLTVLGAKVSQITAGELVDADN 297
 Sbjct: 241 WHLNYQDVIAIGKLFTTGELFTDRIISLAGPQVKNPRLVTRRLGANLSQLTANELNAGEN 300

25 Query: 298 RVISGSVLNGAITQGAHDYLGRYHNQISVIEEGRSKELFGWVAPQPDKYSITRTTLGHFL 357
 Sbjct: 301 RVISGSVLGATAAGPVDYLGRYALQVSVLAEGREKELFGWIMPGSDKFSITRTVLGHFG 360

30 Query: 358 KNKLFKFTTAVNGGDRAMVPIGTIERVMXXXXXXXXXXXXXXXXXVGDTSAXXXXXXXXXX 417
 Sbjct: 361 K-KLFNFTTAVHGGGERAMVPIGAYERVMPLDIPTLLRLDLAAGDTSAXNLGCLELDEE 419

Query: 418 XXXXSFCVCPGKYEXGPLLRKVLETXEKEG 447
 ++VCPGK GP+LR LE EKEG

ORF22ng-1 also shows homology with the OMP from *A.pleuropneumoniae*:

35 gi|1185395 (U24492) 48 kDa outer membrane protein [Actinobacillus
 pleuropneumoniae] Length = 449
 Score = 555 bits (1414), Expect = e-157
 Identities = 284/450 (63%), Positives = 337/450 (74%), Gaps = 4/450 (0%)

40 Query: 27 MIKIKKGLNLPIAGRPEQVIYDGPVITEVALLGEEYVGMPSMKIKEGEAVKKGQVLFED 86
 Sbjct: 1 MITIKKGLDLPIAGTPAQVIHNGNTVNEVAMLGEEYVGMPSMKVREGDVVKKGQVLFED 60

45 Query: 87 KKNPGVVFTAPASGKIAAIHRGEKRVLSVVI AVEGNDEIEFERYVPEALAKLSSEKVR 146
 Sbjct: 61 KKNPGVVFTAPASGTVVTINRGEKRVLSVVIKVEGDEQITFTRYEAAQLASLSAEQVKQ 120

50 Query: 147 NLIQSGLWLTALRTRPFSKIPAVDAEPFAIFVNAMDTNPLAADPTVIIKEAAEDFKRGLLV 206
 Sbjct: 121 NLIESGLWTAFRTRPFSKVPALDAIPSSIFVNAMDTNPLAADPEVVLKEYETDFKDGLTV 180

55 Query: 207 LSRL--TERKIHVCKAAGADVP--SENAANIETHEFGGPHAGLSGTHIHFIIEPVGANKTV 263
 Sbjct: 181 LTRLFNGQKPVYLCCKDADSNIPSPATEGITIKSFSGVHPAGLVGTHIHFDVDPGATKQV 240

60 Query: 264 WTINYQDVIAIGRLFVTGRNLNTERVVALGGLQVKNPRLRLTVLGAKVSQITAGELVDADN 323
 Sbjct: 241 WHLNYQDVIAIGKLFTTGELFTDRIISLAGPQVKNPRLVTRRLGANLSQLTANELNAGEN 300

65 Query: 324 RVISGSVLNGAIAQGAHDYLGRYHNQISVIEEGRSKELFGWVAPQPDKYSITRTTLGHFL 383
 Sbjct: 301 RVISGSVLGATAAGPVDYLGRYALQVSVLAEGREKELFGWIMPGSDKFSITRTVLGHFG 360

Query: 384 KNKLFKFTTAVNGGDRAMVPIGTIERVMXXXXXXXXXXXXXXXXXVGDTSAXXXXXXXXXX 443
 Sbjct: 361 K-KLFNFTTAVHGGGERAMVPIGAYERVMPLDIPTLLRLDLAAGDTSAXNLGCLELDEE 419

Query: 444 XXXXSFCVCPGKYEYGPLLRKVLETIEKEG 473
 ++VCPGK YGP+LR LE IEKEG

70 Sbjct: 420 DLALCTYVCPGKNNYGPMRLRAALEKIEKEG 449

Based on this analysis, including the homology with the outer membrane protein of *Actinobacillus pleuropneumoniae*, it was predicted that these proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

ORF22-1 (35.4kDa) was cloned in pET and pGex vectors and expressed in *E.coli*, as described above. The products of protein expression and purification were analyzed by SDS-PAGE. Figure 5A shows the results of affinity purification of the GST-fusion protein, and Figure 5B shows the results of expression of the His-fusion in *E.coli*. Purified GST-fusion protein was used to immunise mice, whose sera were used for ELISA (positive result) and FACS analysis (Figure 5C). These experiments confirm that ORF22-1 is a surface-exposed protein, and that it is a useful immunogen.

10 Example 16

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 133>:

```

1   .GCGnCGnAAA TCATCCATCC CC..nACGTC GTAGGCCCTG AAGCCAACTG
51  GTTTTTTATG GTAGCCAGTA CGTTTGTGAT TGCTTTGATT GGTATTTTGG
101 TTAAGTAAAA AATCGTCGAA CCGCAATTGG GCCCTTATCA ATCAGATTTG
151 TCACAAGAAG AAAAAGACAT TCGGCATTCC AATGAAATCA CGCCTTTGGA
201 ATATAAAGGA TTAATTGGG CTGGCGTGGT GTTTGTTGCC TTATCCGCCC
251 TATTGGCTTG GAGCATCGTC CCTGCCGACG GTATTTTGCG TCATCCTGAA
301 ACAGGATTGG TTTCCGGTTC GCCGTTTTTA AAATCGATTG TTGTTTTTAT
351 TTTCTTGTTG TTTGCACTGC CGGGCATTGT TTATGGCCGG GTAACCCGAA
20 401 GTTTCGCGCG CGAACAGGAA GTCGTTAATG CGmyGGCCGA ATCGATGAGT
451 ACTCTGGsGC TTTmTTGsw CAkcATCTTT TTTGCCGCAC AGTTTGTGCG
501 ATTTTTTAAT TGGACGAATA TTGGGCAATA TATTGCCGTT AAAGGGGCGA
551 CGTTCTTAAA AGAAGTCGGC TTGGGCGGCA GCGTGTTGTT TATCGGTTTT
601 ATTTTAATTT GTGCTTTTAT CAATCTGATG ATAGGCTCCG CCTCCGCGCA
25 651 ATGGGCGGTA ACTGCCCGCA TTTTCGTCCC TATGCTGATG TTGGCCGGCT
701 ACGCGCCCGA AGTCATTCAA GCCGCTTACC GCATCGGTGA TTCCGTTACC
751 AATATTATTA CGCCGATGAT GAGTTATTTT GGGCTGATTA TGGCGACGGT
801 GrkCmmTAC AAAAAGATG CGGGCGTGGG TaCGcTGATT wCTATGATGT
851 TGCCGTATTC CGCTTTCTTC TTGATTGCGt GGATTGCCTT ATTCTGCATT
30 901 TGGGTATTTg TTTTGGGCCT GCCCGTCGGT CCCGGCGCGC CCACATTCTA
951 TCCCGCACCT TAA

```

This corresponds to the amino acid sequence <SEQ ID 134; ORF12>:

```

1   .AXXIIHPXXV VGPEANWFFM VASTFVIALI GYFVTEKIVE PQLGPYQSDL
51  SQEEKDIRHS NEITPLEYKG LIWAGVVFVA LSALLAWSIV PADGILRHPE
35 101 TGLVSGSPFL KSIVVFIFLL FALPGIVYGR VTRSLRGEQE VVNAXAESMS
151 TLXLXLXXIF FAAQVFAFFN WTNIGQYIAV KGATFLKEVG LGGSVLFIGF
201 ILICAFINLM IGSASAQWAV TAPIFVPLM LAGYAPEVIQ AAYRIGDSVT
251 NIITPMSYF GLIMATVXXY KKDAGVGTLI XMMLPYSAFF LIAWIALFCI
301 WVFVLGLPVG PGAPTFYPAP *

```

40 Further sequence analysis revealed the complete DNA sequence <SEQ ID 135> to be:

```

1   ATGAGTCAAA CCGATACGCA ACGGACGGA CGATTTTAC GCACAGTCGA
51  ATGGCTGGGC AATATGTTGC CGCATCCGGT TACGCTTTT ATTATTTTCA
101 TTGTGTTATT GCTGATTGCC TCTGCCGTCG GTGCGTATT CGGACTATCC
151 GTCCCCGATC CGCGCCCTGT TGGTGCGAAA GGACGTGCCG ATGACGGTPT
45 201 GATTACATT GTCAGCTGC TCAATGCCGA CGGTTTTATC AAAATCCTGA
251 CGCATACCGT TAAAAATTTT ACCGGTTTCG CGCCGTTGGG AACGGTGTTG
301 GTTCTTTTAT TGGCGTGGG GATTGCGGAA AAATCGGGCT TGATTTCCGC
351 ATTAATGCGC TTATTGCTCA CAAAATCGCC ACGCAAATC ACTACTTTTA
401 TGTTTGTTTT TACAGGGATT TTATCTAATA CCGCTTCTGA ATGGGCTAT
50 451 GTCGTCCTAA TCCCTTTGTC CGCCATCATC TTTCATTCCC TCGGCCGCCA
501 TCCGCTTGCC GGTCTGGCTG CGGCTTTCGC CGGCGTTTCG GCGGTTATT

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551 CGGCCAATCT GTTCTTAGGC ACAATCGATC CGCTCTTGGC AGGCATCACC
 601 CAACAGGCGG CGCAAATCAT CCATCCCGAC TACGTCGTAG GCCCTGAAGC
 651 CAACTGGTTT TTTATGGTAG CCAGTACGTT TGTGATTGCT TTGATGGTT
 701 ATTTTGTAC TGAAAAATC GTCGAACCGC AATTGGGCCC TTATCAATCA
 751 GATTTGTAC AAGAAGAAAA AGACATTCGG CATTCCAATG AAATCACGCC
 801 TTTGGAATAT AAAGGATTAA TTTGGGCTGG CGTGGTGTTC GTTGCCATTAT
 851 CCGCCCTATT GGCTTGGAGC ATCGTCCCTG CCGACGGTAT TTTGCGTCAT
 901 CCTGAAACAG GATTGGTTTC CGGTTGCGCG TTTTAAAT CGATTGTTGT
 951 TTTTATTTTC TTGTTGTTTG CACTGCCGGG CATTGTTTAT GGCCGGGTAA
 1001 CCCGAAGTTT GCGCGCGGAA CAGGAAGTCG TTAATGCGAT GGCCGAATCG
 1051 ATGAGTACTC TGGGGCTTTA TTTGGTCATC ATCTTTTGTG CCGCACAGTT
 1101 TGTCGCATTT TTTAATTGGA CGAATATTGG GCAATATATT GCCGTTAAAG
 1151 GGGCGACGTT CTTAAAGAA GTCCGGCTGG GCGGCAGCGT GTTGTTTATC
 1201 GGTTTTATTT TAATTGTGTC TTTTATCAAT CTGATGATAG GCTCCGCCCTC
 1251 CCGCAATGG GCGGTAAGTC CGCCGATTTT CGTCCCTATG CTGATGTTGG
 1301 CCGGTACGC GCCCGAAGTC ATTCAAGCCG CTTACCGCAT CGGTGATTCC
 1351 TTACCAATA TTATTACGCC GATGATGAGT TATTTGCGGC TGATTATGGC
 1401 GACGGTGATC AAATACAAAA AAGATGCGGG CGTGGGTACG CTGATTCTTA
 1451 TGATGTTGCC GTATTCGCTC TTCTTCTTGA TTGCGTGGAT TGCCTTATTC
 1501 TGCATTGGG TATTTGTTT GGGCCTGCC GTCCGTCCCG GCGCGCCAC
 1551 ATTCTATCCC GCACCTTAA

This corresponds to the amino acid sequence <SEQ ID 136; ORF12-1>:

1 MSQTDTRDQ RFLRTVEWLG NMLPHPVTLF IIFIVLLLIA SAVGAYFGLS
 51 VFDPRPVGAK GRADDGLIYI VSLLNADGFI KILTHTVKNF TGFAPLGTVL
 101 VSLLCVGIAE KSGGLISALMR LLLTKSPRKL TTFMVVETGI LSNTASELGY
 151 VVLIPLSAII FHSLSGRHPLA GLAAAFAGVS GGYSANLFLG TIDPLLAGIT
 201 QQAQIIHPD YVVGPEANWF FMVASTFVIA LIGYFVTEKI VEPQLGPYQS
 251 DLSQEEKDIR HSNEITPLEY KGLIWAGVVF VALSALLAWS IVPADGILRH
 301 PETGLVSGSP FLKSIVVFIF LLEFALPGIVY GRVTRSLRGE QEVVNMAES
 351 MSTLGLYLV IFFAAQFVAF FNWTNIGQYI AVKGATFLKE VGLGGSVLF
 401 GFILICAFIN LMIGSASAQW AVTAPIFVPM LMLAGYAPEV IQAAYRIGDS
 451 VTNIITPMMS YFGLIMATVI KYKKDAGVGT LISMMLPYSA FFLIAWIALF
 501 CIWVFLGLP VGPGATFYFP AP*

Computer analysis of this amino acid sequence gave the following results:

35 Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF12 shows 96.3% identity over a 320aa overlap with an ORF (ORF12a) from strain A of *N. meningitidis*:

				10	20	30
40	orf12.pep			AXXIIHPXXVVGPEANWFFMVASTFVIALI		
	orf12a	AAAFAGVSGGYSANLFLGTIDPLLAGITQQAQIIHPDYVVGPEANWFFMVASTFVIALI				
		180 190 200 210 220 230				
45	orf12.pep	GYFVTEKIVEPQLGPYQSDLSQEEKDIRHSNEITPLEYKGLIWAGVVFVALSALLAWSIV				
	orf12a	GYFVTEKIVEPQLGPYQSDLSQEEKDIRHSNEITPLEYKGLIWAGVVFVALSALLAWSIV				
		240 250 260 270 280 290				
50	orf12.pep	PADGILRHPETGLVSGSPFLKSIVVFIFLLFALPGIVYGRVTRSLRGEQEVVNAXAESMS				
	orf12a	PADGILRHPETGLVSGSPFLKSIVVFIFLLFALPGIVYGRVTRSLRGEQEVVNAXAESMS				
		300 310 320 330 340 350				
55	orf12.pep	TLXLXLIIFFAAQFVAFFNWTNIGQYIAVKGATFLKEVGLGGSVLFIFILICAFINLM				
	orf12a	TLGLYLVIIFFAAQFVAFFNWTNIGQYIAVKGATFLKEVGLGGSVLFIFILICAFINLM				
60		360 370 380 390 400 410				
	orf12.pep	IGSASAQWAVTAPIFVPMMLAGYAPEVIQAAYRIGDSVTNIITPMMSYFGLIMATVXXY				
		220 230 240 250 260 270				

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orf12a	IGSASAQWAVTAPIFVPMMLLAGYAPEVIQAAYRIGDSVTNIITPMMSYFGLIMATVIKY
	420 430 440 450 460 470
5	orf12.pep
	280 290 300 310 320
	KKDAGVGTLLIXMMLPYSAFFLIWIALFCIWVFLGLPVGPGAPTFYPAPX
	orf12a
	KKDAGVGTLLISMMLPYSAFFLIWIALFCIWVFLGLPVGPGAPTFYPAPX
	480 490 500 510 520

The complete length ORF12a nucleotide sequence <SEQ ID 137> is:

10	1	ATGAGTCAAA	CCGATACGCA	ACGGGACGGA	CGATTTTAC	GCACAGTCGA
	51	ATGGCTGGGC	AATATGTTGC	CGCACCCGCT	TACGCTTTT	ATTATTTTCA
	101	TTGTGTTATT	GCTGATTGCC	TCTGCCGCCG	GTGCGTATT	CGGACTATCC
	151	GTCCCCGATC	CGCGCCCTGT	TGGTGCAGAA	GGACGTGCCG	ATGACGGTTT
	201	GATTCACGTT	GTCAGCCTGC	TGCATGCTGA	CGGTTTGATC	AAAATCCTGA
15	251	CGCATACCGT	TAAAAATTTT	ACCGGTTTCG	CGCCGTTGGG	AACGGTGTTG
	301	GTTTCTTTAT	TGGGCGTGGG	GATTGCGGAA	AAATCGGGCT	TGATTTCCGC
	351	ATTAATGCGC	TTATTGCTCA	CAAAATCTCC	ACGCAAACTC	ACTACTTTTA
	401	TGGTTGTTTT	TACAGGGATT	TTATCTAATA	CCGCTTCTGA	ATTGGGCTAT
	451	GTCGTCTTAA	TCCCTTTGTC	CGCCATCATC	TTTCATTCCC	TCCGCCGCCA
20	501	TCCGCTTGCC	GGTCTGGCTG	CGGCTTTCGC	CGGCGTTTCG	GGCGGTTATT
	551	CGGCCAATCT	GTTCTTAGGC	ACAATCGATC	CGCTCTTGCC	AGGCATCACC
	601	CAACAGGCGG	CGCAAATCAT	CCATCCCGAC	TACGTCGTAG	GCCCTGAAGC
	651	CAACTGGTTT	TTTATGGTAG	CCAGTACGTT	TGTGATTGCT	TTGATTGGTT
	701	ATTTTGTTAC	TGAAAAAATC	GTCGAACCGC	AATTGGGCCC	TTATCAATCA
25	751	GATTGTGCAC	AAGAAGAAAA	AGACATTCCA	CATTCCAATG	AAATCACGCC
	801	TTTGAATAT	AAAGGATTAA	TTTGGGCTGG	CGTGGTGTTT	GTTGCCCTAT
	851	CCGCCCTATT	GGCTTGGAGC	ATCGTCCCTG	CCGACGGTAT	TTTGCGTCAT
	901	CCTGAAACAG	GATTGGTTTC	CGGTTCGCCG	TTTTTAAAT	CAATTGTTGT
	951	TTTTATTTTC	TTGTTGTTTG	CACTGCCGGG	CATTGTTTAT	GGCCGGGTAA
30	1001	CCCGAAGTTT	GCGCGGCGAA	CAGGAAGTCG	TTAATGCGAT	GGCCGAATCG
	1051	ATGAGTACTC	TGGGGCTTTA	TTTGGTCATC	ATCTTTTTTG	CCGCACAGTT
	1101	TGTCGCATTT	TTTAATTGGA	CGAATATTGG	GCAATATATT	GCCGTAAAG
	1151	GGGCGACGTT	CTTAAAGAA	GTCGGCTTGG	GCGGCAGCGT	GTTGTTTATC
	1201	GGTTTTATTT	TAATTGTGTC	TTTATCAAT	CTGATGATAG	GCTCCGCCTC
35	1251	CGCGCAATGG	GCGGTAACAG	CGCCGATTTT	CGTCCCTATG	CTGATGTTGG
	1301	CCGGCTACGC	GCCCGAAGTC	ATTCAAGCCG	CTTACCGCAT	CGGTGATTCC
	1351	GTTACCAATA	TTATTACGCC	GATGATGAGT	TATTTCCGGC	TGATTATGGC
	1401	GACGGTGATC	AAATACAAAA	AAGATGCGGG	CGTGGGTACG	CTGATTTCTA
	1451	TGATGTTGCC	GTATTCGCT	TTCTTCTGA	TTGCGTGGAT	TGCCTTATTC
40	1501	TGCATTTGGG	TATTTGTTTT	GGGCTGCCC	GTCGCTCCG	GCGGCCCCAC
	1551	ATTCTATCCC	GCACCTTAA			

This encodes a protein having amino acid sequence <SEQ ID 138>:

	1	MSQTDTRQDGR	RFLRTVEWLG	NMLPHPVTLF	IIFIVLLLIA	SAAGAYFGLS
	51	VDPDRPVGAK	GRADDGLIHV	VSLLDADGLI	KILTHTVKNF	TGFAPLGTVL
45	101	VSLLGVGIAE	KSLGISALMR	LLLTSPKRL	TTFMVFTGI	LSNTASELGY
	151	VVLIPLSAII	FHSLGRHPLA	GLAAAFAGVS	GGYSANLFLG	TIDPLLAGIT
	201	QQAQIIHPD	YVVGPEANWF	FMVASTFVIA	LIGYFVTEKI	VEPQLGPYQS
	251	DLSQEEKDIR	HSNEITPLEY	KGLIWAGVVF	VALSALLAWS	IVPADGILRH
	301	PETGLVSGSP	FLKSIVVFIF	LLFALPGIVY	GRVTRSLRGE	QEVVNAMAES
50	351	MSTLGLYLVI	IFFAAQFVAF	FNWNTNIGQYI	AVKGATFLKE	VGLGGSVLFI
	401	GFILICAFIN	LMIGSASQW	AVTAPIFVPM	LMLAGYAPEV	IQAAAYRIGDS
	451	VTNIITPMMS	YFGLIMATVI	KYKKGAVGT	LISMMLPYSA	FFLIAWIALF
	501	CIWVFLGLP	VGPGAPTFYP	AP*		

55 ORF12a and ORF12-1 show 99.0% identity in 522 aa overlap:

		10	20	30	40	50	60
	orf12a.pep	MSQTDTRQDGR	RFLRTVEWLG	NMLPHPVTLF	IIFIVLLLIA	SAAGAYFGLS	VDPDRPVGAK
60	orf12-1	MSQTDTRQDGR	RFLRTVEWLG	NMLPHPVTLF	IIFIVLLLIA	SAVGAAYFGLS	VDPDRPVGAK
		10	20	30	40	50	60
		70	80	90	100	110	120
	orf12a.pep	GRADDGLIHV	VSLLDADGLI	KILTHTVKNF	TGFAPLGTVL	VSLLGVGIAE	KSLGISALMR
65	orf12-1	GRADDGLIY	VSLLDADGFI	KILTHTVKNF	TGFAPLGTVL	VSLLGVGIAE	KSLGISALMR

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		70	80	90	100	110	120
		130	140	150	160	170	180
5	orf12a.pep	LLLTSPRKLTTFMVVFTGILSNTASELGYVVLIPLSAIFHSLGRHPLAGLAAAFAGVS					
	orf12-1	LLLTSPRKLTTFMVVFTGILSNTASELGYVVLIPLSAIFHSLGRHPLAGLAAAFAGVS					
		130	140	150	160	170	180
		190	200	210	220	230	240
10	orf12a.pep	GGYSANLFLGTIDPLLAGITQQAQIIHPDYVVGPEANWFFMVASTFVIALIGYFVTEKI					
	orf12-1	GGYSANLFLGTIDPLLAGITQQAQIIHPDYVVGPEANWFFMVASTFVIALIGYFVTEKI					
		190	200	210	220	230	240
		250	260	270	280	290	300
15	orf12a.pep	VEPQLGPFYQSDLSQEEKDIRHSNEITPLEYKGLIWAGVVFVALSALLAWSIVPADGILRH					
	orf12-1	VEPQLGPFYQSDLSQEEKDIRHSNEITPLEYKGLIWAGVVFVALSALLAWSIVPADGILRH					
		250	260	270	280	290	300
		310	320	330	340	350	360
	orf12a.pep	PETGLVSGSPFLKSIVVFIFLLFALPGIVYGRVTRSLRGEQEVVNAMAE SMSTLGLYLVI					
	orf12-1	PETGLVSGSPFLKSIVVFIFLLFALPGIVYGRVTRSLRGEQEVVNAMAE SMSTLGLYLVI					
25		310	320	330	340	350	360
		370	380	390	400	410	420
	orf12a.pep	IFFAAQFVAFFNWTNIGQYIAVKGATFLKEVGLGGSVLFIFGILICAFINLMIGSASAQW					
30	orf12-1	IFFAAQFVAFFNWTNIGQYIAVKGATFLKEVGLGGSVLFIFGILICAFINLMIGSASAQW					
		370	380	390	400	410	420
		430	440	450	460	470	480
	orf12a.pep	AVTAPIFVPMMLLAGYAPEVIQAAYRIGDSVTNIITPMSYFGLIMATVIKYKKGAGVGT					
35	orf12-1	AVTAPIFVPMMLLAGYAPEVIQAAYRIGDSVTNIITPMSYFGLIMATVIKYKKGAGVGT					
		430	440	450	460	470	480
		490	500	510	520		
40	orf12a.pep	LISMMLPYSAFFLIWIALFCIWVFLGLFVGPAGPTFYFAPX					
	orf12-1	LISMMLPYSAFFLIWIALFCIWVFLGLFVGPAGPTFYFAPX					
		490	500	510	520		

45 Homology with a predicted ORF from *N.gonorrhoeae*

ORF12 shows 92.5% identity over a 320aa overlap with a predicted ORF (ORF12.ng) from *N. gonorrhoeae*:

	orf12.pep	AXXIHPXXVVGPEANWFFMVASTFVIALI	30
50	orf12ng	AAAFAGVSGGYSANLFLGTIDPLLAGITQQAQIIHPDYVVGPEANWFFMAASTFVIALI	232
	orf12.pep	GYFVTEKIVEPQLGPFYQSDLSQEEKDIRHSNEITPLEYKGLIWAGVVFVALSALLAWSIV	90
	orf12ng	GYFVTEKIVEPQLGPFYQSDLSQEEKDIRHSNEITPLEYKGLIWAGVVFVALSALLAWSIV	292
55	orf12.pep	PADGILRHPETGLVSGSPFLKSIVVFIFLLFALPGIVYGRVTRSLRGEQEVVNAXAESMS	150
	orf12ng	PADGILRHPETGLVAGSPFLKSIVVFIFLLFALPGIVYGRITRSLRGEREVVNAXAESMS	352
60	orf12.pep	TLXLXLXIIFFAAQFVAFFNWTNIGQYIAVKGATFLKEVGLGGSVLFIFGILICAFINLM	210
	orf12ng	TLGLYLVIIFFAAQFVAFFNWTNIGQYIAVKGAVFLKKFRLGGSVLFIFGILICAFINLM	412
	orf12.pep	IGSASQWAVTAPIFVPMMLLAGYAPEVIQAAYRIGDSVTNIITPMSYFGLIMATVXXY	270
65	orf12ng	IGSASQWAVTAPIFVPMMLLAGNAPQVIQAAYRIGDSVTNIITPMSYFGLIMATVIKY	472

5	1	ATGAGTCAAA	CCGACGCGCG	TCGTAGCGGA	CGATTTTAC	GCACAGTCGA
	51	ATGGCTGGGC	AATATGTTGC	CGCACCCGGT	TACGCTTTTT	ATTATTTTCA
	101	TTGTGTATT	GCTGATTGcc	tctgCCGTCG	GTGCGTATT	CGGACTATCC
	151	GTCCCCGATC	CGCGTCTGT	TGGGGCGAAA	GGAGCTGCG	ATGACGGTT
	201	GATTACAGTT	GTCAGCTGC	TCGATGCCGA	CGGTTTGATC	AAAATCCTGA
10	251	CGCATACCGT	TAAAAATTTT	ACCGGTTTCG	CGCGTITGGG	AACGGTGT
	301	GTTTCTTTAT	TGGGCGTGGG	GATTGCGGAA	AAATCGGGCT	TGATTTCCGC
	351	ATTAATGCGC	TTATTGCTCA	CAAAATCCCC	ACGCAAACTC	ACTACTTTTA
	401	TGGTTGTTTT	TACAGGGATT	TTATCCAATA	CGGCTTCTGA	ATTGGGCTAT
	451	GTCGTCTTAA	TCCCTTTGTC	CGCCGTATC	TTTCATTTCG	TCGGCCGCCA
15	501	TCCGCTTGGC	GGTTTGGCTG	CGGCTTTCGC	CGGCGTTTTCG	GGCGGTTATT
	551	CGGCCAATCT	GTTCTTAGGC	ACAATCGATC	CGCTCTTGGC	AGGCATCACC
	601	CAACAGGCGG	GTCAAATCAT	CCATCCCGAC	TACGTCGTAG	GCCCTGAAGC
	651	CAACTGGTTT	TTTATGGCAG	CCAGTACGTT	TGTGATTGCT	TTGATTGGTT
	701	ATTTTGTATC	TGAAAAATC	GTCGAACCCG	AATTGGGCCC	TTATCAATCA
20	751	GATTTGTCAC	AGAAAGAAAA	AGACATTTCG	CATTCCAATG	AATCATGCGC
	801	TTTGGAATAT	AAAGGATTAA	TTTGGGCAGG	CGTGGTGTTT	GTTGCCCTAT
	851	CCGCCCTATT	GGCTTGAGAG	ATCGTCCCTG	CCGACCGTAT	TTTGCGTCAT
	901	CCTGAAACAG	GATTGGTTGC	CGGTTCCGCG	TTTTTAAAA	CGATTGTGTG
	951	TTTTATTTTC	TGTGTTTGTG	CGCTGCCGGG	CATGTTTAT	GGCCGGATAA
25	1001	CCCGAAGTTT	GCGCGGCGAA	CGGGAAGTCG	TTAATGCGAT	GGCCGAATCG
	1051	ATGAGTACTT	TGGGACTTTA	TTTGGTCATC	ATCTTTTTTG	CCGCACAGTT
	1101	TGTCGCATTT	TTTAATTGGA	CGAATATTGC	GCAATATATT	GCGGTTAAAG
	1151	GGGCGGTGTT	CTTAAAAGAA	GTCGGCTTGG	GCGGCAGTGT	TGTGTTTATC
	1201	GGTTTTATTT	TAATTTGTGC	TTTTATCAAT	CTGATGATAG	GCTCCGCCCTC
30	1251	CGCGCAATGG	GCGGTAAGTC	CGCCGATTTT	CGTCCCTATG	CTGATGTTGG
	1301	CCGGCTACGC	GCCCGAAGTC	ATTCAAGCCG	CTTACCGCAT	CGGTGATTC
	1351	GTTACCAATA	TTATTACGCC	GATGATGAGT	TATTTCCGGC	TGATTATGGC
	1401	GACGGTAATC	AAATACAAAA	AAGATGCGGG	CGTAGGCACG	CTGATTTCTA
	1451	TGATGTTGCC	GTATTCGCGT	TTCTTCTTAA	TTGCATGGAT	CGCCTTATTC
35	1501	TGCATTTGGG	TATTTGTTTT	GGGCTGCCCC	GTCGGTCCCC	GCACACCCAC
	1551	ATTCTATCCG	GTGCTTAA			

	1	MSQTDARRSG	RFLRTVEWLG	NMLPHPVTLF	IIFIVLLLIA	SAVGAYFGLS
40	51	VPDPRPVGAK	GRADDGLIHV	<u>VSLLDADGLI</u>	KILTHTVKNF	TGFAPLGTVI
	101	<u>VSLLGVGIAE</u>	KSGLISALMR	LLLTKSPRKL	TTFMVVFTGI	LSNTASELGY
	151	<u>VVLIPLSAVI</u>	FVSLGRHPLA	GLAAAFAGVS	GGYSANFLFG	TIDPLLAGIT
	201	<u>QQAQIHPD</u>	VYVVGPEANWF	FMAASTFVIA	LIGYFVTEKI	VEPQLGPYQS
	251	DLSEQEEDIR	HSNEITPLEY	<u>KGLIWAGVVF</u>	<u>VALSALLAWS</u>	IVPADGILRH
45	301	PETGLVAGSP	FLKSIVVFIF	<u>LLFALPGIVY</u>	GRITRSLRGE	REVVNAMAES
	351	<u>MSTLGLYLVI</u>	<u>IFFAAQFVAF</u>	<u>FNWNTNIGQYI</u>	AVKGAVFLKK	FRLGGSVLFI
	401	GFILICAFIN	LMIGSASAQW	AVTAPIFVPM	LMLAGNAPQV	IQAAYRIGDS
	451	<u>VTNIITPMMS</u>	<u>YFGLIMATVI</u>	KYKKDAGVGT	LISMMLPYSA	<u>FLIAWIALF</u>
	501	CIWVFVLGLP	VGPPTPTFPY	VP*		

50		10	20	30	40	50	60
	orf12-1.pep	MSQTD	TQRDGRFLRTVEW	LGNNMLPHPV	TLFIIFIV	LLLIASAVG	AYFGLSVDP
			:::				
	orf12ng	MSQTD	ARRSGRFLRTVEW	LGNNMLPHPV	TLFIIFIV	LLLIASAVG	AYFGLSVDP
		10	20	30	40	50	60
55		70	80	90	100	110	120
	orf12-1.pep	GRADD	GLIYIVSLLNADG	FIKIL	THTVKNFTG	FAPLGT	VLVSLLGVG
			:::				
	orf12ng	GRADD	GLIHVVSLLDADG	LILKIL	THTVKNFTG	FAPLGT	VLVSLLGVG
		70	80	90	100	110	120
60		130	140	150	160	170	180
	orf12-1.pep	LLLT	KSPRKL	TTFMV	VFTGIL	SNTASE	LGYYVLI
65	orf12ng	LLLT	KSPRKL	TTFMV	VFTGIL	SNTASE	LGYYVLI

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		130	140	150	160	170	180
		190	200	210	220	230	240
5	orf12-1.pep	GGYSANLFLGTIDPLLAGITQQAQIIHPDYVVGPEANWFFMVASTFVIALIGYFVTEKI					
	orf12ng	GGYSANLFLGTIDPLLAGITQQAQIIHPDYVVGPEANWFFMAASTFVIALIGYFVTEKI					
		190	200	210	220	230	240
		250	260	270	280	290	300
10	orf12-1.pep	VEPQLGPYQSDLSQEEKDIRHSNEITPLEYKGLIWAGVVFVALSALLAWSIVPADGILRH					
	orf12ng	VEPQLGPYQSDLSQEEKDIRHSNEITPLEYKGLIWAGVVFVALSALLAWSIVPADGILRH					
		250	260	270	280	290	300
		310	320	330	340	350	360
15	orf12-1.pep	PETGLVSGSPFLKSIVVFIFLLFALPGIVYGRVTRSLRGEQEVVNAMAESMSTLGLYLVI					
	orf12ng	PETGLVAGSPFLKSIVVFIFLLFALPGIVYGRITRSLRGEREVVNAMAESMSTLGLYLVI					
		310	320	330	340	350	360
		370	380	390	400	410	420
	orf12-1.pep	IFFAAQFVAFFNWTNIGQYIAVKGATFLKEVGLGGSVLFIFILICAFINLMIGSASAQW					
	orf12ng	IFFAAQFVAFFNWTNIGQYIAVKGAVFLKEVGLGGSVLFIFILICAFINLMIGSASAQW					
25		370	380	390	400	410	420
		430	440	450	460	470	480
	orf12-1.pep	AVTAPIFVPMMLAGYAPEVIQAAAYRIGDSVTNIITPMMSYFGLIMATVIKYKKDAGVGT					
30	orf12ng	AVTAPIFVPMMLAGYAPEVIQAAAYRIGDSVTNIITPMMSYFGLIMATVIKYKKDAGVGT					
		430	440	450	460	470	480
		490	500	510	520		
35	orf12-1.pep	LISMMLPYSAFFLIAWIALFCIWFVFLGGLPVGPAPTFFYPAPX					
	orf12ng	LISMMLPYSAFFLIAWIALFCIWFVFLGGLPVGPPTFFYPVPX					
		490	500	510	520		

In addition, ORF12ng shows significant homology with a hypothetical protein from *E.coli*:

40	sp P46133 YDAH_ECOLI HYPOTHETICAL 55.1 KD PROTEIN IN OGT-DBPA INTERGENIC REGION >gi 1787597 (AE000231) hypothetical protein in ogt 5' region [Escherichia coli] Length = 510 Score = 329 bits (835), Expect = 2e-89 Identities = 178/507 (35%), Positives = 281/507 (55%), Gaps = 15/507 (2%)	
45	Query: 8 RSGRFLRTVEWLGNNMLPHPVTTXXXXXXXXXXASAVGAYFGLSVDPDRPVGAKGRADDGL 67 +SG+ VE +GN +PHP +A+ + FG+S +P D Sbjct: 13 QSGKLYGWVERIGNKVPHPFLLEFYLIIVLMVTTAILSAFGVSAKNP-----TDGTP 64	
50	Query: 68 IHVVSLLDADGLIKILHTHTVKNFTGFAPXXXXXXXXXXIAEKSGLSALMRLLLTKSP 127 + V +LL +GL L + +KNF+GFAP +AE+ GL+ ALM + + Sbjct: 65 VVVKNNLSVEGLHWFLPNVIKNFSGFAPLGAILALVLGAGLAERVGLLPALMVKMASHVN 124	
55	Query: 128 RKLTTFMVVFVGILSNTASELGYVVLIPLSAVIFHSLGRHPLAGLAAAFAGVSGGYSANL 187 + ++MV+F S+ +S+ V++ P+ A+IF ++GRHP+AGL AA AGV G++ANL Sbjct: 125 ARYASYMVLFIAFFSHISSDAALVIMPPMGALIFLAVGRHPVAGLLAAIAGVCGGFTANL 184	
60	Query: 188 FLGTIDPLLAGITQQAQIIHPDYVVGPEANWFFMAASTFVIALIGYFVTEKIVEPQLGP 247 + T D LL+GI+ +AA +P V NW+FMA+S V+ ++G +T+KI+EP+LG Sbjct: 185 LIVTTDVLVLSGISTEAAAAFNQMHVSVIDNWFMASSVVVLTIVGGLTIDKIIIEPRLGQ 244	
65	Query: 248 YQSDLSQEEKDIRHSNEITPLEYKGLIWAGVVFVALSALLAWSIVPADGILRHPETGLVA 307 +Q + ++ + + S GL AGVV + A +A ++P +GILR P V Sbjct: 245 WQGNSEKLQITITESQRF-----GLRIAGVVSLLFIAAIALMVIPQNGILRDPINHTVM 298	
70	Query: 308 GSFFLKSIIVVFIFLLFALPGIVYGRITRSLRGEREVVNAMAESMSTLGLYLXXXXXXX 367 SPF+K IV I L F + + YG TR++R + ++ + M E M + ++ Sbjct: 299 PSFFIKGIVPLIILFFVVSLEYGIATRTIRROADLPHLMIEPMKEMAGFIVMVFLPAQF 358	
	Query: 368 XXXXNWTNIGQYIAVKGAVFLKEVGLGGSVLFIFILICAFINLMIGSASAQWAVTAPIF 427 NW+N+G++IAV L+ GL G F+G L+ +F+ + I S SA W++ APIF	

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Sbjct: 359 VAMFNWSNMGKFIAGVGLTDILESSGLSGIPAFVGLALLSSFLCMFIASGSASISILAPIF 418
 Query: 428 VPMLMLAGYAPEVIQAAAYRIGDSVTNIITPMMSYFGLIMATVIKYKKDAGVGTLSMMLP 487
 VPM ML G+ P Q +RI DS + P+ + L + + +YK DA +GT S++LP
 Sbjct: 419 VPMFMLLGFHFAFAQILFRIADSSVLPLAPVSPFVPLFLGFLQRYKPDAKLGTYYSLVLP 478
 Query: 488 YSAFFLIAWIALFCIWVFLVGLPVGPG 514
 Y FL+ W+ + W +++GLP+GPG
 Sbjct: 479 YPLIFLVVWLLMLLAW-YLVGLPIGPG 504

Based on this analysis, including the presence of several putative transmembrane domains and the predicted actinin-type actin-binding domain signature (shown in bold) in the gonococcal protein, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 17

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 141>:

1 ..ACAGCCGGCG CAGCAGGTTn CnCGGTCTTC GTTTTCGTAA CGGACAGTCA
 51 GGTGGAGGTG TTCGGAACA TCCAGACCGC AGTGGAACA GGTTTTTTTC
 101 ATGGCATTTC GGTTCGTCT GTGTTGGTG CGGCGGCACA AGACTCGGCA
 151 ATgGCTTCGC GCAGTGCCTC TATACCGGTA TTTTCAGCAA CGGAAATGCG
 201 GACGGcGgCA ATTTTCCCG CAGCGTCGCG CCATATGCCC GTGTTTgTT
 251 CTTCAGACGG CAGCAGGTCG GTTTGTGTGT ACACCTTgAT GCACGGAaTA
 301 TCGCCGGCAT GGATTCTTG CAGTACGTTT TCCACGTCTT CAATCTGCTG
 351 TCCGCTGTTC GGAGCGGCGG CATCGACGAC GTGCAGCAGC ACATCgGcTT
 401 gCGCGGTTTC TTCCAGCGTG GCgGAAAGG CGGAAATCAG TTTgTGCGGC
 451 agATyGCTnA CGAATCCGAC GGTATCGGTC AGGATAATGC TGCATTCGGG
 501 ACT..

This corresponds to the amino acid sequence <SEQ ID 142; ORF14>:

1 ..TAGAAGXXVF VEVTDSEQVEV FGNIQTAVET GFFHGISVSS VFGAAQDSA
 51 MASRSASIPV FSATEMRTAA IFPAASRHMP VFCSSDGSRS VLLYTLMHGI
 101 SPAWISCSTF STSSICCPLE GAAASTTCSS TSACAVSSSV AEKAEISLCG
 151 RXLTNPTVSV RIMLHSG..

Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF14 shows 94.0% identity over a 167aa overlap with an ORF (ORF14a) from strain A of *N.meningitidis*:

				10	20	30
orf14.pep				TAGAAGXXVF	VEVTDSEQVEV	FGNIQTAVET
				:	:	: : :
orf14a	GRQLGFLRVGGALFVITAQARVNNALCDCLTTGAAGFAV	FVFDGQM	QVFGNVQPAVET			
	150	160	170	180	190	200
		40	50	60	70	80
orf14.pep	GFFHGISVSSV	FVGAAQDSAMASRSASIPVFSATEMRTAAIFPAASRHMPVFCSSDGSRS				
orf14a	GFFHGISVSSV	FVGAAQYSAMASRSASIPVFSATEMRTAAIFPAASRHMPVFCSSDGSRS				
	210	220	230	240	250	260
		100	110	120	130	140
orf14.pep	VLLYTLMHGISPAWISCSTFSTSSICCPLEFGAAASTTCSST	SACAVSSSV	AEKAEISLCG			
orf14a	VLLYTLMHGISPAWISCSTFSTSSICCPLEFGAAASTTCSST	SACAVSSSV	AEKAEISLCG			
	270	280	290	300	310	320

160
 orf14.pep RXLTNPTVSVRIMLHSG
 | | | | | | | | | |
 orf14a RSLTNPTVSVRIMLHSGGLMYSRRRAVVSSVAKSWSFAYMPDLVSRNLRLDLPTLVX
 330 340 350 360 370 380

The complete length ORF14a nucleotide sequence <SEQ ID 143> is:

	1	ATGGAGGATT	TGCAGGAAAT	CGGGTTCGAT	GTCCGCCGCC	TAAAGGTAGG
	51	TCGGCAGCGC	GAACATCATC	GTCTGCATCA	TCCCCAGCCC	GGCACAAGCGG
10	101	AGGCGGACGA	TGTATTGTTT	CGCTTCTTTT	TGTTTGGCGG	CTTCGATTTT
	151	TTGCGCGTCA	TAGGGTCCGG	CGGTGTAGCC	TATCTGCCTG	ATTTTCAACA
	201	GAATGTCGGA	AAGGCGGATT	TTGCCGTCGT	CCCAGACGAC	GCGGCAGCGG
	251	TGCGTGTCTG	AATTGAGGTC	GATGCGGACG	ATGCCGTCTG	TACGCAAAAAG
	301	CTGCTGTTCG	ATCAGCCAGA	CGCAGGCGCG	CGAGGTGATG	CGCCCGAGCA
15	351	TTAAAACCGC	CTCGCGCGTG	CCGCCGTGGG	TTTCCACAAA	GTCGGACTGG
	401	ACTTCGGGCA	GGTCGTACAG	GCGGATTGGG	TCGAGGATTT	CTTGGGGCGG
	451	CAGCTCGGTT	TTTTGCGCGT	CGGCCGTGCG	TGTGTTGTAA	TAACTGCCCC
	501	AGCCCGCGTC	ATAATGCTTT	TGTGCGACTG	CCTGACAACC	GCGCGCAGAG
	551	GTTTCGCGGT	CTTCGTTTTT	GTAACGGACG	GTCAGATGCA	GGTTTTTCGG
20	601	AACGTCACAG	CCGCAGTGGA	AACAGGTTTT	TTTCATGGCA	TTTCGGTTTT
	651	GTCTGTGTTT	GGTGCAGCGG	CACAATACTC	GGCAATGGCT	TCCGCGCAGT
	701	CGCTCTATAC	GTCATTTTCA	GCAACGGAAA	TGCGGACGCG	GGCAATTTTT
	751	CCCGCAGCGT	CGCGCCATAT	GCCCGTGTTT	TGTTCTTTAG	ACGGCAGCAG
	801	GTCCGGTTTTG	TTGTACACCT	TGATGCACGG	AATATCGCCG	GCATGATATT
25	851	CTTGCACTAC	GTTTCCACG	TCTTCAATCT	GCTGTCCGCT	GTTCCGGAGG
	901	CGGCGATCGA	CGACGTGCAG	CAGCACATCG	GCTTGGCGGG	TTTCTTCCAG
	951	CGTGGCGGAA	AAGGCGGAAA	TCAGTTTGTG	CGGCAGATCG	CTGACGAATC
	1001	CGACGGTATC	GGTCAGGATA	ATGCTGCATT	CGGGACATGAT	GTACAGCCGC
	1051	CGCGCCGTTC	TGTCGAGTGT	GGCGAAAAGC	TGGTCTTTTC	CATATATGCC
	1101	CGACTTGGTC	AGCCGGTTGA	ACAGACTGGA	TTTGGCGACA	TTGGTATAG

30 This encodes a protein having amino acid sequence <SEQ ID 144>:

35

1	MEDLQEIGFD	VAAVKVGRQR	EHHRLHHPQP	GNGEADDVLF	AFFLVGGGDF
51	LRVIGCGGVA	YLPDFQQNVG	KADFAVVPDD	AAAVRAVIEV	DADDAVCTQK
101	LLFDQPDAGQ	AGDAAEH*NR	LARAAVGFHK	VGLDFGQVVQ	ADLVDFELGR
151	QLGFLRVGGA	LFVITQAQRV	NNALCDCLTT	GAAGFVAVFV	VTDGQMQVFG
201	NVQPAVETGF	FHGISVSSVF	GAAQYSAMA	SRSASIPVFS	ATEMRTAAIF
251	PAASRHPVTF	CSSDGSRSVL	LYTLMHGIST	AWISCTSFST	SSICCPLEGA
301	AASTTCSSTS	ACAVSSSVAE	KAEISLCGRS	LTNPTVSVRI	MLHSGLMYSR
351	RAVVSSVAKS	WSFAYMPDLV	SRLNRDLPT	LV*	

It should be noted that this sequence includes a stop codon at position 118.

40 Homology with a predicted ORF from *N.gonorrhoeae*

ORF14 shows 89.8% identity over a 167aa overlap with a predicted ORF (ORF14.ng) from *N. gonorrhoeae*:

	orf14.pep	TAGAAGXXVFVFVTDTSQVEVFGNIQTAVET	30
		: : : : : : : :	
45	orf14.ng	GRQGFGRVVGASFVITAQAGIDDALCDCLTADAAGFAVFAFVADGQMQVFGNVQPAVET	208
	orf14.pep	GFFHGISVSSVFGAAAQDSAMASRSASIPVFSATEMRTAAIFPAASRHPVFCSSDGSR	90
50	orf14.ng	GFFHGISVSSVFGAAQYSAMASRSASIPVFSATEMRTAAIFPAASRHPVFCSSDGSR	268
	orf14.pep	VLLYTLMHGISPAWISCSTFTSTSSICPLFGAAASTTCSSTSACAVSSSVAEKAEISLCG	150
	orf14.ng	VLLYTLMHGISWAWISCSTFTSTSSICPLFRAAASTTCSSTSACTVSSKVAEKAEISLCG	328
55	orf14.pep	RXLTNPTVSVRIMLHSG	167
	orf14.ng	RSLTNPTVSVRIMLHAGLMYSRRVAVSRVAKSWSFAYMPDLVSRNLRLDPTLV	382

The complete length ORF14ng nucleotide sequence <SEQ ID 145> is predicted to encode a protein having amino acid sequence <SEO ID 146>:

```

      1 MEDLQEIGFD VAAVKVGRQR EHHRLHHTQS GNGKADDVLF AFFLVGGFDF
    51 LRVIGCGGVA CLPDFQQNVG EADFAVVPDD AAARAVIEV DADDAVCAQK
   101 LLFDQPDAGG AGNAAEHQHC FVRAIMGFHK VGLDFGQVVQ ADLVEDFLGR
   151 QFGFFRVGGA SFVITAQAGI DDALCDCLTA DAAGFAVFAP VADGQMVFVG
    201 NVQPAVETGF FHGISVSSVF GAAQYSAMA SRSASIPVFS ATEMRTAAIF
   251 PAASRHMPVF CSSDGSRSVL LYTLMHGISW AWISCSTFST SSICCPLFRA
   301 AASTTCSSTS ACTVSSKVAE KAEISLCGRS LTNPTVSVRI MLHAGLMYSR
   351 RAVVSRVAKS WSFAYMPDLV SRLNRLDLPT LV*

```

Based on the putative transmembrane domain in the gonococcal protein, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 18

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 147>:

```

    1 ..GGCCATTACT CCGACCGCAC TTGGAAGCCG CGTTTGNGCG GCCGCCGTCT
   51 GCCGTATCTG CTTTATGGCA CGCTGATTGC GGTATTGTG ATGATTTTGA
  101 TGCCGAAGTC GGGCAGCTTC GGTTCGGCT ATGCGTCGCT GCGCGCTTTG
   151 TCGTTCGGCG CGCTGATGAT TCGCTGTGTA GACGTGTCGT CAAATATGGC
   201 GATGCAGCCG TTTAAGATGA TGGTCGGCGA CATGGTCAAC GAGGAGCAGA
   251 AAA.NTACGC CTACGGGATT CAAAGTTTCT TAGCAAATAC GGGCGCGGTC
   301 GTGGCGGCGA TTCTGCCGTT TGTGTTTGCG TATATCGGTT TGGCGAACAC
   351 CGCCGANAAA GGCGTTGTGC CGCAGACCGT GGTCTGTGGC TTTTATGTGG
   401 GTGCGGCGTT GCTGGTGATT ACCAGCGCGT TCACGATTTT CAAAGTGAAG
   451 GAATACGANC CGGAAACCTA CGCCCGTTAC CACGGCATCG ATGTCGCCGC
   501 GAATCAGGAA AAGCCAACT GGATCGCACT CTTAAAA.CC GCGC..

```

This corresponds to the amino acid sequence <SEQ ID 148; ORF16>:

```

    1 ..GHYSDRTWKP RLXGRRLPYL LYGTLIAVIV MILMPNSGSF GFGYASLAAL
   51 SFGALMIALL DVSSNMAMQP FKMMVGDVMN EEQKXYAYGI QSFLANTGAV
  101 VAAILPFVFA YIGLANTAXK GVVPQTVVVA FVGAALLVI TSAFTIFKVK
  151 EYXPETYARY HGIDVAANQE KANWIALLKX A..

```

Further work revealed the complete nucleotide sequence <SEQ ID 149>:

```

    1 ATGTCGGAAT ATACGCCTCA AACAGCAAAA CAAGGTTTGC CCGCGCTGGC
   51 AAAAAGCAGC ATTTGGATGC TCAGTTTCGG CTTTCTCGGC GTTCAGACGG
  101 CCTTTACCTT GCAAAGCTCG CAAATGAGCC GCATTTTTCA AACGCTAGGC
   151 GCAGACCCGC ACAATTGGG CTGGTTTTTC ATCCTGCCGC CGCTGGCGGG
   201 GATGCTGGTG CAGCCGATTG TCGGCCATTA CTCGACCCG ACTTGAAGC
   251 CCGGTTTGGG CGGCGCCGT CTGCCGATC TGCTTTATGG CACGCTGATT
   301 GCGGTTATTG TGATGATTTT GATGCCGAAC TCGGGCAGCT TCGGTTTCGG
   351 CTATGCGTCG CTGGCGGCTT TGTGTTTCGG CGCGCTGATG ATTGCGCTGT
   401 TAGACGTGTC GTCAAATATG GCGATGCAGC CGTTTAAGAT GATGGTCGGC
   451 GACATGGTCA ACGAGGAGCA GAAAGGCTAC GCCTACGGGA TTCAAAGTTT
   501 CTTAGCAAAT ACGGGCGCGG TCGTGGCGGC GATTCTGCCG TTTGTGTTTG
   551 CGTATATCGG TTTGGCGAAC ACCGCCGAGA AAGGCGTTGT GCCGCAGAC
   601 GTGGTCGTGG CGTTTATGT GGGTGCGCGG TTGCTGGTGA TTACCAGCGC
   651 GTTCACGATT TTCAAAGTGA AGGAATACGA TCCGGAACCC TACGCCCGTT
   701 ACCACGGCAT CGATGTCGCC GCGAATCAGG AAAAAGCCAA CTGGATCGAA
   751 CTTTGAAAAA CCGCGCCTAA GCGTTTTTGG ACGGTTACTT TGGTGCAATT
   801 CTTCTGCTGG TTCGCCTTCC AATATATGTG GACTTACTCG GCAGGCGCGA
   851 TTGCGGAAAA CGTCTGGCAC ACCACCGATG CGTCTCCGT AGGTATCAG
   901 GAGGCGGGTA ACTGGTACGG CGTTTGGCG GCGGTGCAGT CGGTTGCGGC
   951 GGTGATTTGT TCGTTGTAT TGGCGAAACT GCCGAATAAA TACCATAAGG
  1001 CGGGTTATTT CCGCTGTTTG GCTTGGGCG CGCTCGGCTT TTTCTCGGTT
  1051 TTCTTCATCG GCAACCAATA CCGCTGGTG TTGCTTATA CCTAATCGG
  1101 CATCGCTTGG GCGGGCATT TCACTTATCC GCTGACGATT GTGACCAACG
  1151 CCTTGTCTGG CAAGCATATG GGCATTACT TGGGCTTGT TAACGGCTCT
  1201 ATCTGTATGC CTCAAATCGT CGCTTCGCTG TTGAGTTTCG TGCTTTTCCC
  1251 TATGCTGGG GCGTTCAGG CCACTATGTT CTTGGTAGG GCGCTCGTCC
  1301 TGCTGCTGGG CGCGTTTTCC GTGTTCTTGA TTAAGAAAC ACACGGCGGG
  1351 GTTTGA

```

This corresponds to the amino acid sequence <SEQ ID 150; ORF16-1>:

```

      1  MSEYTPQTAK  QGLPALAKST  IWMLSFGLG  VQTAFTLQSS  QMSRIFQTLG
     51  ADPHNLGWFF  ILPLAGMLV  QPIVGHYSR  TWKPRLGRR  LPYLLYGTLI
    101  AVIVMILMPN  SGSFGFGYAS  LAALSFGALM  IALLDVSSNM  AMQPFKMMVG
15    151  DMVNEEQKGY  AYGIQSFLAN  TGAVVAAILP  FVFAYIGLAN  TAEKGVPQT
     201  VVVAFYVGAA  LLVITSaftI  FKVKEYDPET  YARYHGIDVA  ANQEKANWIE
     251  LLKTAPKAFW  TVTLVQFFCW  FAFQYMWYTS  AGAIAENVWH  TTDASSVGyQ
     301  EAGNWYGVLA  AVQSVAAVIC  SFVLAKVPNK  YHKAGYFGCL  ALGALGFFSV
     351  FFIGNQYALV  LSYTLIGIAW  AGIITYPLTI  VTNALSGKHM  GTYLGLFNgs
10    401  ICMPQIVASL  LSFVLFPMLG  GLQATMFLVG  GVVLLLGAFS  VFLIKETHGG
     451  V*

```

Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF16 shows 96.7% identity over a 181aa overlap with an ORF (ORF16a) from strain A of *N.*

15 *meningitidis*:

```

                                     10      20      30
orfl6.pep                        GHYSDRTWKPRXLXGRRLPYLLYGTLIAVIV
                                     |||
20  orfl6a      IFQTLGADPHSLGWFFILPLPLAGMLVQPIVGHYSDRTWKPRLGRRLPYLLYGTLIAVIV
                   50      60      70      80      90     100

                                     40      50      60      70      80      90
orfl6.pep      MILMPNSGSFGFGYASLAALSFGALMIALLDVSSNMAMQPFKMMVGDMVNEEQXYAYGI
                                     |||
25  orfl6a      MILMPNSGSFGFGYASLAALSFGALMIALLDVSSNMAMQPFKMMVGDMVNEEQKYAYGI
                   110     120     130     140     150     160

                                     100     110     120     130     140     150
orfl6.pep      QSFLANTGAVVAAILPFVFAYIGLANTAXKGVPQTVVVAFYVGAALLVITSaftIFKVK
                                     |||
30  orfl6a      QSFLANTGAVVAAILPFVFAYIGLANTAEGKVPQTVVVAFYVGAALLVITSaftIFKVK
                   170     180     190     200     210     220

                                     160     170     180
orfl6.pep      EYXPETYARYHGIDVAANQEKANWIALLKXA
                                     ||
35  orfl6a      EYNPETYARYHGIDVAANQEKANWIELLKTA PKAFWTVTLVQFFCWFAFQYMWYTSAGAI
                   230     240     250     260     270     280

40  orfl6a      AENVWHTTDASSVGyQEAGNWYGVLA AVQSVAAVICSFVLAKVPNKYHKAGYFGCLALGA
                   290     300     310     320     330     340

```

The complete length ORF16a nucleotide sequence <SEQ ID 151> is:

```

      1  ATGTCGGAAT  ATACGCCTCA  AACAGCAAAA  CAAGGTTTGC  CCGCGCTGGC
     51  AAAAAGCACG  ATTTGGATGC  TCAGTTTCGG  CTTTCTCGGC  GTTCAGACGG
45    101  CCTTTACCCT  GCAAAGCTCG  CAGATGAGCC  GCATCTTCCA  GACGCTCGGT
     151  GCCGATCCGC  ACAGCCTCGG  CTGGTTCTTT  ATCCTGCCGC  CGCTGGCGGG
     201  GATGCTGGTG  CAGCCGATTG  TCGGCCATTA  CTCCGACCGC  ACTTGGGAAGC
     251  CGCGTTTGGG  CGGCCGCCGT  CTGCCGTATC  TGCTTTATGG  CACGCTGATT
     301  GCGGTTATTG  TGATGATTTT  GATGCCGAAC  TCGGGCAGCT  TCGGTTTCGG
50    351  CTATGCGTCG  CTGGCGGCTT  TGTCTGTCGG  CGCGCTGATG  ATTGCGCTGT
     401  TAGACGTGTC  GTCAAATATG  GCGATGCAGC  CGTTTAAGAT  GATGGTCGGC
     451  GACATGGTCA  ACGAGGAGCA  GAAAGGCTAC  GCCTACGGGA  TTCAAAGTTT
     501  CTTAGCGAAT  ACGGGCGCGG  TCGTGCGCGC  GATTCTGCCG  TTTGTGTTTG
55    551  CGTATATCGG  TTTGGCGAAC  ACCGCCGAGA  AAGGCGTTGT  GCCGCAGACC
     601  GTGGTCGTGG  CGTTTATGTG  GGGTGCGGCG  TTGCTGGTGA  TTACCAGCGC
     651  GTTCACGATT  TTCAAAGTGA  AGGAATACAA  TCCGGAACCC  TACGCCCGTT
     701  ACCACGGCAT  CGATGTCGCC  GCGAATCAGG  AAAAAGCCAA  CTGGATCGAA
     751  CTCTTGAAAA  CCGCGCCTAA  GGCGTTTGGG  ACGGTTACTT  TGGTGCAATT
     801  CTTCTGCTGG  TTCGCCTTCC  AATATATGTG  GACTTACTCG  GCAGGCGCGA
60    851  TTGCGGAAAA  CGTCTGGCAC  ACCACCGATG  CGTCTTCCGT  AGGTTATCAG
     901  GAGGCGGGTA  ACTGGTACGG  CGTTTGGGCG  GCGGTGCAGT  CGGTTGCGGC
     951  GGTGATTGTG  TCGTTTGTAT  TGGCGAAAGT  GCCGAATAAA  TACCATAAGG

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5
 1001 CGGGTTATTT CGGCTGTTTG GCTTTGGGCG CGCTCGGCTT TTTCTCCGTT
 1051 TTCTTCATCG GCAACCAATA CGCGCTGGTG TTGTCTTATA CCTTAATCGG
 1101 CATCGCTTGG GCGGGCATT TCACTTATCC GCTGACGATT GTGACCAACG
 1151 COTTGTCCGG CAAGCATATG GGCACCTACT TGGGCCTGTT TAACGGCTCT
 1201 ATCTGTATGC CGCAAATCGT CGCTTCGCTG TTGAGTTTCG TGCTTTTCCC
 1251 TATGCTGGGC GGCTTGCAGG CCACTATGTT CTTGGTAGGG GGCGTCGTCC
 1301 TGCTGCTGGG CGCGTTTTCG GTGTTCTGA TTAAAGAAAC ACACGGCGGG
 1351 GTTTGA

This encodes a protein having amino acid sequence <SEQ ID 152>:

10
 1 MSEYTPQTAK QGLPALAKST IWMLSFGLG VQTAFTLQSS QMSRIFQTLG
 51 ADPHSLGWFF ILPPLAGMLV QPIVGHYS DR TWKPRLGRR LPYLLYGTLI
 101 AVIVMILMPN SGSFGFGYAS LAALSFGALM IALLDVSSNM AMQPFKMMVG
 151 DMVNEEQGY AYGIQSFLAN TGAVVAAILP FVFAYIGLAN TAEKGVVPQT
 201 VVVAFYVGAA LLVITSFTI FKVKEYNPET YARYHGIDVA ANQEKANWIE
 15
 251 LLKTAPKAFW TVTLVQFFCW FAFQYMWYS AGAIAENVWH TTDASSVGYQ
 301 EAGNWyGVLA AVQSVAVIC SFVLAKVPNK YHKAGYFGCL ALGALGFFSV
 351 FFIGNQYALV LSYTLIGIAW AGIITYPLTI VTNALSGKHM GTYLGLEFNGS
 401 ICMFQIVASL LSFVLFPM LG GLQATMFLVG GVVLLLGAFS VFLIKETHGG
 451 V*

20 ORF16a and ORF16-1 show 99.6% identity in 451 aa overlap:

		10	20	30	40	50	60
orfl6a.pep		MSEYTPQTAKQGLPALAKSTIWMLSFGLGVQTAFTLQSSQMSRIFQTLGADPHSLGWFF					
orfl6-1		MSEYTPQTAKQGLPALAKSTIWMLSFGLGVQTAFTLQSSQMSRIFQTLGADPHNLGWFF					
		10	20	30	40	50	60
orfl6a.pep		70	80	90	100	110	120
orfl6-1		ILPPLAGMLVQPIVGHYS DRTWKPRLGRR LPYLLYGTLIAVIVMILMPNSGSFGFGYAS					
		70	80	90	100	110	120
orfl6a.pep		130	140	150	160	170	180
orfl6-1		LAALSFGALMIALLDVSSNMAMQPFKMMVGDMDVNEEQKGYAYGIQSFLANTGAVVAAILP					
		130	140	150	160	170	180
orfl6a.pep		190	200	210	220	230	240
orfl6-1		FVFAYIGLANTA EKGVPQT VVVAFYVG AALLVITSFTIFKVKEYNPETYARYHGIDVA					
		190	200	210	220	230	240
orfl6a.pep		250	260	270	280	290	300
orfl6-1		ANQEKANWIELLKTAPKAFWTVTLVQFFCWFAFQYMWYSAGAIAENVWHTTDASSVGYQ					
		250	260	270	280	290	300
orfl6a.pep		310	320	330	340	350	360
orfl6-1		EAGNWyGVLA AVQSVAVIC SFVLAKVPNK YHKAGYFGCLALGALGFFSVFFIGNQYALV					
		310	320	330	340	350	360
orfl6a.pep		370	380	390	400	410	420
orfl6-1		LSYTLIGIAWAGIITYPLTI VTNALSGKHMGT YLGLEFNGSICMPQIVASLLSFVLFPM LG					
		370	380	390	400	410	420
orfl6a.pep		430	440	450			
orfl6-1		GLQATMFLVGGVVLLLGAFSVFLIKETHGGVX					
		430	440	450			

Homology with a predicted ORF from *N.gonorrhoeae*

ORF16 shows 93.9% identity over a 181aa overlap with a predicted ORF (ORF16.ng) from *N. gonorrhoeae*:

5	orf16.pep	GHYSDRTWKPRXLGRRLLPYLLYGTLLIAVIV	30
	orf16ng	HFSNARRRPAQFGLVFHPAAAGGDAGSADSGYYSRTWKPRLGRRLLPYLLYGTLLIAVIV	131
10	orf16.pep	MILMPNSGSFGFGYASLAALSFGALMIALLDVSSNMAMQPFKMMVGDVNEEQKXYAYGI	90
	orf16ng	MILMPNSGSFGFGYASLAALSFGALMIALLDVSSNMAMQPFKMMVGDVNEEQKXYAYGI	191
15	orf16.pep	QSFLANTGAVVAAILPFVFAYIGLANTAXKGVVPQTVVVAFYVGAALLVITSFTIFKVK	150
	orf16ng	QSFLANTDAVVAAILPFVFAYIGLANTAEGVVPQTVVVAFYVGAALLIITSFTISKVK	251
	orf16.pep	EYXPETYARYHGIDVAANQEKANWIALLKXA	181
	orf16ng	EYDPETYARYHGIDVAANQEKANWFELLKTAPKVFVTVTPVQFFCWFAPRYMWTYSAGAI	311

20 The complete length ORF16ng nucleotide sequence <SEQ ID 153> is:

1	ATGATAGGGG	ATCGCCGCGC	CGGCAACCAT	TTCGGATTTT	CCAAAGCAAA
51	TACTTTTCAA	ATCAAAAAAA	AGGATTTACT	TTATGTCGGA	ATATACGCCT
101	CAAACAGCAA	AACAAGGTTT	GCCCGCGCCG	GCAAAAAGCA	CGATTGGAT
151	GTTGAGCTTC	GGCTATCTCG	GCGTTCAGAC	GGCCTTTACC	CTGCAAAGCT
201	CGCAGATGAG	CCGCATTTTT	CAAACGCTAG	GCGCAGACCC	GCACAATTTG
251	GGCTGGTTTT	TCATCCTGCC	GCCGCTGGCG	GGGATGCTGG	TTACGCCGAT
301	AGTGGCTACT	ACTCAGACCG	CACTTGGAAG	CCGCGCTTGG	GCGGCCGCCG
351	CCTGCCGTAT	CTGCTTTACG	GCACGCTGAT	TGCGGTCTATC	GTGATGATTT
401	TGATGCCGAA	CTCGGGCAGC	TTGCGTTTCG	GCTATGCGTC	GCTGGCGGCC
451	TTGTGCTTCG	GCGCGCTGAT	GATTGCGCTG	TTGGACGTGT	CGTCGAATAT
501	GGCGATGCAG	CCGTTTAAGA	TGATGGTCCG	CGATATGGTC	AACGAGGAGC
551	AGAAAAGCTA	CGCCTACGGG	ATTCAAAGTT	TCTTAGCGAA	TACGGACGCG
601	GTTGTGGCAG	CGATTCTGCC	GTTTGTGTTT	GCGTATATCG	GTTTGGCGAA
651	CACTGCCGAG	AAAGGCGTTG	TGCCACAAAC	CGTGGTCGTA	GCATTCTATG
701	TGGGTGCGGC	GTTACTGATT	ATTACCAAGT	CGTTCACAA	CTCCAAAGTC
751	AAAGAAATAC	ACCCGGAAC	CTACGCCCGT	TACCACGGCA	TCGATGTGCG
801	CGCGAATCAG	GAAAAAGCCA	ACTGGTTCGA	ACTCTTAAAA	ACCGCGCCTA
851	AAGTGTTCG	GACGCTTACT	CCGGTACAGT	TTTTCTGCTG	GTTTCGCTTC
901	CGGTATATGT	GGACTTACTC	GGCAGGCGCG	ATTGCAGAAA	ACGTCTGGCA
951	CACTACCGAT	GCGTCTTCCG	TAGGCCATCA	GGAGGCGGGC	AACCGGTACG
1001	GCGTTTGGC	GGCGGTGTAG			

This encodes a protein having amino acid sequence <SEQ ID 154>:

1	MIGDRRAGNH	FGFSKANTFQ	IKKKDLLYVG	IYASNSKTRF	ARAGKKHDL
51	VELRLSRRSD	GLYPAKLADE	PHFSNARRRP	AQFGLVFHPA	AAGGDAGSAD
101	SGYYSRTWK	PRLGGRRLPY	LLYGTLLIAVI	VMILMPNSGS	FGFGYASLAA
151	LSFGALMIAL	LDVSSNMAMQ	PFKMMVGDV	NEEQKXYAYG	IQSFLANTDA
201	VVAAILPFVF	AYIGLANTAE	KGVVPQTVV	AFYVGAALLI	ITSFTISKV
251	KEYDPETYAR	YHGIDVAANQ	EKANWFELLK	TAPKVFVTVT	PVQFFCWFAP
301	RYMWTYSAGA	IAENVHHTTD	ASSVGHQEAG	NRYGVLAAV*	

50 ORF16ng and ORF16-1 show 89.3% identity in 261 aa overlap:

		30	40	50	60	70	80
	orf16-1.pep	MLSFGFLGVQTAFTLQSSQMSRIFQTLGADPHNLGWFFILPPLAGMLVQPI-VGHYSDRT					
	orf16ng	DVELRLSRRSDGLYPAKLADEPHFSNARRRPAQFGLVF-HPAAAGGDAGSADSGYYSRT					
55		50	60	70	80	90	100
	orf16-1.pep	WKPRLGRRLLPYLLYGTLLIAVIVMILMPNSGSFGFGYASLAALSFGALMIALLDVSSNMA					
	orf16ng	WKPRLGRRLLPYLLYGTLLIAVIVMILMPNSGSFGFGYASLAALSFGALMIALLDVSSNMA					
60		110	120	130	140	150	160

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		150	160	170	180	190	200
5	orf16-1.pep	MQPFKMMVGD MNVEEQKGYAYGIQSFLANTGAVVAAILPFVFAYIGLANTAEGVVPQTV					
	orf16ng	MQPFKMMVGD MNVEEQKSYAYGIQSFLANTDAVVAAILPFVFAYIGLANTAEGVVPQTV					
		170	180	190	200	210	220
10	orf16-1.pep	VVAFYVGAALLVITS AFTIFKVKEYDPETYARYHGIDVAANQEKANWIELLK TAPKAFWT					
	orf16ng	VVAFYVGAALLIITS AFTISKVKEYDPETYARYHGIDVAANQEKANWFELLK TAPKVFWT					
		230	240	250	260	270	280
15	orf16-1.pep	VTLVQFFCWF AFRYMW TYSAGAIAENVWHTTDASSVGYQEAGN WYGVLA AVQSVA A VICS					
	orf16ng	VTPVQFFCWF AFRYMW TYSAGAIAENVWHTTDASSVGHQEAGN RYGVLA AVX					
		290	300	310	320	330	340

- 20 Based on this analysis, including the presence of several putative transmembrane domains in the gonococcal protein, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 19

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 155>:

25	1	ATGTTGTTCC	GTAAACGAC	CGCCGCCGTT	TGGCGCATA	CCTTGATGCT
	51	GAACGGCTGT	ACGTTGATGT	TGTGGGGAAT	GAACAACCCG	GTCAGCGAAA
	101	CAATCACCCG	NAAACACGTT	GNCAAAGACC	AAATCCGNGN	CTTCGGTGTG
	151	GTTGCCGAAG	ACAATGCCCA	ATTGGAAAAG	GGCAGCCTGG	TGATGATGGG
	201	CGGAAAATAC	TGGTTCGTCG	TCAATCCCGA	AGATTCGGCG	AA.NTGACGG
30	251	GNATTTTGAN	GGCAGGGCTG	GACAAACCCCT	TCCAAATAGT	TNAGGATACC
	301	CCGAGCTATG	C.TGCCACCA	AGCCCTGCCG	GTCAAACCTCG	GATCGNCTGG
	351	CAGCCAGAAT...				

This corresponds to the amino acid sequence <SEQ ID 156; ORF28>:

35	1	MLFRKTTAAV	LAHTLMLNGC	TLMLWGMNPN	VSETITRKHV	XKDQIRXFGV
	51	VAEDNAQLEK	GSLVMMGGKY	WFFVNPEDSA	XTTGILXAGL	DKPFQIVXDT
	101	PSYXCHQALP	VKLGSXGSQN...			

Further work revealed the complete nucleotide sequence <SEQ ID 157>:

	1	ATGTTGTTCC	GTAAACGAC	CGCCGCCGTT	TGGCGGCAA	CCTTGATGCT
	51	GAACGGCTGT	ACGTTGATGT	TGTGGGGAAT	GAACAACCCG	GTCAGCGAAA
40	101	CAATCACCCG	CAAACACGTT	GACAAAGACC	AAATCCGCGC	CTTCGGTGTG
	151	GTTGCCGAAG	ACAATGCCCA	ATTGGAAAAG	GGCAGCCTGG	TGATGATGGG
	201	CGGAAAATAC	TGGTTCGTCG	TCAATCCCGA	AGATTCGGCG	AAGCTGACGG
	251	GCATTTTGAA	GGCAGGGCTG	GACAAACCCCT	TCCAAATAGT	TGAGGATACC
	301	CCGAGCTATG	CTCGCCACCA	AGCCCTGCCG	GTCAAACCTCG	AATCGCCTGG
45	351	CAGCCAGAAT	TTCAGTACCG	AAGGCCTTTG	CCTGCGCTAC	GATACCGACA
	401	AGCCTGCCGA	CATCGCCCAAG	CTGAAACAGC	TCGGGTTTGA	AGCGGTCAAA
	451	CTCGACAATC	GGACCATTTA	CACGCGCTGC	GTATCCGCCA	AAGGCAAATA
	501	CTACGCCACA	CCGCAAAAAC	TGAACGCCGA	TTACCATTTT	GAGCAAAGTG
50	551	TGCCTGCCGA	TATTTATTAC	ACGGTTACTG	AAGAACATAC	CGACAAATCC
	601	AAGCTGTTTG	CAAATATCTT	ATATACGCCC	CCCTTTTGA	TACTGGATGC
	651	GGCGGGCGCG	GTA CTGGCCT	TGCCTGCGGC	GGCTCTGGGT	GCGGTCGTGG
	701	ATGCCGCCCG	CAATGA			

This corresponds to the amino acid sequence <SEQ ID 158; ORF28-1>:

55	1	MLFRKTTAAV	LAATLMLNGC	TLMLWGMNPN	VSETITRKHV	DKDQIRAFGV
	51	VAEDNAQLEK	GSLVMMGGKY	WFFVNPEDSA	KLTGILKAGL	DKPFQIVEDT
	101	PSYARHQALP	VKLESPGSQN	FSTEGLCRLY	DTDKPADIAK	LKQLGFPAVK
	151	LDNRTIYTRC	VSAKGKYYAT	PQKLNADYHF	EQSVPADIYY	TVTEEHTDKS

201 KLFANILYTP PFLILDAAGA VLALPAAALG AVVDAARK*

Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF28 shows 79.2% identity over a 120aa overlap with an ORF (ORF28a) from strain A of *N.*

5 *meningitidis*:

		10	20	30	40	50	60
orf28.pep		MLFRKTTAAVLAHTLMLNGCTLMWGMNPNVSETITRKHVXKDQIRXFGVVAEDNAQLEK					
		: : : : : : :					
orf28a		MLFRKTTAAVLAATLMLNGCTVMMWGMNSPFSETTARKHVDKDQIRAFGVVAEDNAQLEK					
		10	20	30	40	50	60
		70	80	90	100	110	120
orf28.pep		GSLVMMGGKYWFVNPEDSAXXTGILXAGLDKPFQIVXDTPSYXCHQALPVKLGSGXSQN					
		: : : : : :					
orf28a		GSLVMMGGKYWFVNPEDSAKLTKILKAGLDKQFQMVPEPNRFA-YQALPVKLESPASQN					
		70	80	90	100	110	
orf28a		FSTEGLCRLRYDTRDPADIATKLQLEFEAVELDNRTIYTRCVSAKGKYYATPQKLNADYHF					
		120	130	140	150	160	170

20 The complete length ORF28a nucleotide sequence <SEQ ID 159> is:

	1	ATGTTGTTCC	GTAAACGAC	CGCCGCCGTT	TTGGCGGCAA	CCTTGATGTT
	51	GAACGGCTGT	ACGGTAATGA	TGTGGGTAT	GAACAGCCCG	TTCAGCGAAA
	101	CGACCGCCCG	CAAACACGTT	GACAAGGACC	AAATCCGCGC	CTTCGGTGTG
25	151	GTTGCCGAAG	ACAATGCCCA	ATTGGAAAAG	GGCAGCCTGG	TGATGATGGG
	201	CGGGAAATAC	TGGTTCGTCG	TCAATCCTGA	AGATTGCGCG	AAGCTGACGG
	251	GCATTTTGAA	GGCCGGGTG	GACAAGCAGT	TTCAAATGGT	TGAGCCCAAC
	301	CCGCGCTTTG	CCTACCAAGC	CCTGCCGCTC	AAACTCGAAT	CGCCCGCCAG
	351	CCAGAATTC	AGTACCGAAG	GCCTTGCCT	GCGCTACGAT	ACCGACAGAC
	401	CTGCCGACAT	CGCCAAGCTG	AAACAGCTTG	AGTTGAAGC	GGTCGAATC
30	451	GACAATCGGA	CCATTACAC	GCGCTGCGTC	TCCGCCAAG	GCAAATACTA
	501	CGCCACACCG	CAAAAACCTGA	ACGCCGATTA	TCATTTTGAG	CAAAGTGTGC
	551	CTGCCGATAT	TTATTACAG	GTTACGAAAA	AACATACCGA	CAAATCCAAG
	601	TTGTTTGAAA	ATATTGCATA	TACGCCACAC	ACGTTGATAC	TGGATGCGGT
	651	GGGCGCGGTG	CTGGCCTTGC	CTGTCGCGGC	GTTGATTGCA	GCCACGAATT
35	701	CCTCAGACAA	ATGA			

This encodes a protein having amino acid sequence <SEQ ID 160>:

	1	MLFRKTTAAV	LAATLMLNGC	TVMWGMNSP	FSETTARKHV	DKDQIRAFGV
	51	VAEDNAQLEK	GSLVMMGGKY	WFVNPEDSA	KLTKILKAGL	DKQFQMVPEP
40	101	PRFAYQALPV	KLESPASQNF	STEGLCRLRYD	TDRPADIATKL	KQLEFEAVEL
	151	DNRTIYTRCV	SAKGKYYATP	QKLNADYHFE	QSVPADIIYT	VTKKHTDKSK
	201	LFENIAYTPT	TLILDAVGAV	LALPVAALIA	ATNSSDK*	

ORF28a and ORF28-1 show 86.1% identity in 238 aa overlap:

		10	20	30	40	50	60
orf28a.pep		MLFRKTTAAVLAATLMLNGCTVMMWGMNSPFSETTARKHVDKDQIRAFGVVAEDNAQLEK					
		: : : : : :					
orf28-1		MLFRKTTAAVLAATLMLNGCTLMWGMNPNVSETITRKHVDKDQIRAFGVVAEDNAQLEK					
		10	20	30	40	50	60
		70	80	90	100	110	119
orf28a.pep		GSLVMMGGKYWFVNPEDSAKLTKILKAGLDKQFQMVPEPNRFA-YQALPVKLESPASQN					
		: : : : : :					
orf28-1		GSLVMMGGKYWFVNPEDSAKLTKILKAGLDKPFQIVXDTPSYARHQALPVKLESPASQN					
		70	80	90	100	110	120
		120	130	140	150	160	179
orf28a.pep		FSTEGLCRLRYDTRDPADIATKLQLEFEAVELDNRTIYTRCVSAKGKYYATPQKLNADYHF					
		: : : : : :					
orf28-1		FSTEGLCRLRYDTRDPADIATKLQLEFEAVELDNRTIYTRCVSAKGKYYATPQKLNADYHF					
		130	140	150	160	170	180

ORF28 shows 84.2% identity over a 120aa overlap with a predicted ORF (ORF28.ng) from *N.*

	orf28.pep	MLFRKTTAAVLAHTLMLNGCTLMLWGMNPNVSETITRKHVXKDQIRXFGVVAEDNAQLEK	60
		:: : : : : :	
	orf28ng	MLFRKTTAAVLAATLILNGCTMMLRGMNPNVSTITRKHVVDKQIRAFGVVAEDNAQLEK	60
15	orf28.pep	GSLVMMGGKGYWFVVPNPEDSAXXTGILXAGLDKPFQIVXDTPSYXCHQALPVKLGSGXGSON	120
		:: : : : : :	
	orf28ng	GSLVMMGGKGYWFAVNPEDSAKLTGLLKAGLDKPFQIVEDTPSYARHOALPVKFEAPGSON	120
		:: : : : : :	

20	1	ATGTTGTTCC	GTA AACGAC	CGCGCCGTT	TTGGCGGCAA	CCTTGATACT
	51	GAACGGGTGT	ACGATGATGT	TGCGGGGGAT	GAACAACCCG	GTCAGCCAAA
	101	CAATCACCCG	CAAAACGTT	GACAAAGACC	AAATCCGCGC	CTTCGGTGTG
	151	GTTGCCGAG	ACAATGCCCA	ATTGGAAAAG	GGCAGCTTGG	TGATGATGGG
25	201	CGGGAAATAC	TGGTTCGCG	TCAATCCCGA	AGATTGCGCG	AAGCTGACGG
	251	GCCTTTTGAA	GGCGGGGTG	GACAAGCCCT	TCCAAATAGT	TGAGGATACC
	301	CCGAGCTATG	CCCGCCACCA	AGCCCTGCCG	GTCAAATTCG	AAGCGCCCAG
	351	CAGCCAGAA	TTCAGTACCG	GAGGTCTTTG	CCTGCGCTAT	GATACCGGCA
30	401	GACCTGACGA	CATCGCCAAG	CTGAAACAGC	TTGAGTTTAA	AGCGGTCAAA
	451	CTCGACAATC	GGACCATTTA	CACGCGCTGC	GTATCCGCCA	AAGGCAAAAT
	501	CTACGCCACG	CCGCAAAAAC	TGAACGCCGA	TTATCATTTT	GAGCAAAAGT
	551	TGCCCCGCCG	TATTTATTAT	ACGGTTACTG	AAAAACATAC	CGACAAATCC
	601	AAGCTGTTTG	GAAATATCTT	ATATACGCCC	CCCTTGTTGA	TATTGGATGC
	651	GGCGGCCGCG	GTGCTGGTCT	TGCCTATGGC	TCTGATTGCA	GCCCGGAATT
	701	CCTCAGACAA	ATGA			

35 1 MLFRKTTAAV LAATLILNGC TMMLRGMNPN VSQTIIRKHV DKDQIRAFGV
51 VAEDNAQLEK GSLVMMGGKY WFAVNPEDSA KLTGLLKAGL DKPFQIVEDT
101 PSYARHQALP VKFEAPGSQN FSTGGGLCLRY DTGRPPDIAK LKQLEFKAHV
151 LDNRTTIYTR PSAKGGYYT PQKLNADYHF EQSVPDIIYY TVTEKHTDKS
201 KLFGNILYTP VLLILDAAA VLVLPMALIA AANSDDK*

		10	20	30	40	50	60
	orf28-1.pep	MLFRKTTAAVLAATLMLNGCTLMLWGMNNPVSETITRKHVDKDQIRAFGVVAEDNAQLEK					
		: : : :					
45	orf28ng	MLFRKTTAAVLAATLIILNGCTMMLRGMNNPVSQTITRKHVDKDQIRAFGVVAEDNAQLEK					
		10	20	30	40	50	60
		70	80	90	100	110	120
	orf28-1.pep	GSLVMMGGKYWFVVNPEDSAKL TGILKAGLDKPFQIVEDTPSYARHQALPVKLES PGSQN					
		: : : :					
50	orf28ng	GSLVMMGGKYWFAVNPEDSAKL TGLLKAGLDKPFQIVEDTPSYARHQALPVKFEPG SQN					
		70	80	90	100	110	120
		130	140	150	160	170	180
	orf28-1.pep	FSTEGLC LRYDTDKPADI AKLKQLGF EAVKLDNR TIYTRCVS AKGKYATP QKLNAD YHF					
		: : : :					
55	orf28ng	FSTGGLC LRYDTGRPDDI AKLKQLEF KAVKLDNR TIYTRCVS AKGKYATP QKLNAD YHF					
		130	140	150	160	170	180
		190	200	210	220	230	239
60	orf28-1.pep	EQSVPADIIYYTVTEEHTDKSKLFANILYT PPFILDAAGAVLALPAALGA VVDAARKX					
		: : : :					
	orf28ng	EQSVPADIIYYTVTEKHTDKSKLFGN ILYT PPLLIDAAA AVLVLPMALIAA ANSSDKX					

190 200 210 220 230

Based on this analysis, including the presence of a putative transmembrane domain in the gonococcal protein, it was predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

ORF28-1 (24kDa) was cloned in pET and pGex vectors and expressed in *E.coli*, as described above. The products of protein expression and purification were analyzed by SDS-PAGE. Figure 6A shows the results of affinity purification of the GST-fusion protein, and Figure 6B shows the results of expression of the His-fusion in *E.coli*. Purified GST-fusion protein was used to immunise mice, whose sera were used for ELISA, which gave a positive result. These experiments confirm that ORF28-1 is a surface-exposed protein, and that it may be a useful immunogen.

Example 20

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 163>:

```

1  .GTCAGTCCTG TACTGCCTAT TACACACGAA CGGACAGGGT TTGAAGGTGT
51  TATCGGTTAT GAAACCCATT TTTCAGGGCA CGGACATGAA GTACACAGTC
101 CGTTCGATCA TCATGATTCA AAAAGCACTT CTGATTTCAG CGGCGGTGTA
151 GACGGCGGTT TTAAGTGTTA CCAACTTCAT CGAACATGGT CGGAAATCCA
201 TCCGGAGGAT GAATATGACG GGCCGCAAGC AGCG.ATTAT CCGCCCCCCC
251 GAGGAGCAAG GGATATATAC AGCTATTATG TCAAAGGAAC TTCAACAAAA
20  301 ACAAAGACTA GTATTGTCCC TCAAGCCCCA TTTTCAGACC GTTGGCTAGA
351 AGAAAATGCC GGTGCCGCCT CTGGT..

```

This corresponds to the amino acid sequence <SEQ ID 164; ORF29>:

```

1  .VSPVLPITHE RTGFEGVIGY ETHFSGHGHE VHSFPDHHDS KTSDFSGGV
51  DGGFTVYQLH RTWSEIHPED EYDGPQAAXY PPEGGARDIY SYYVKGTSTK
25  101 TKTSIVPQAP FSDRWLEENA GAASG..

```

Further work revealed the complete nucleotide sequence <SEQ ID 165>:

```

1  ATGAATTTGC CTATTCAAAA ATTCATGATG CTGTTTGCAG CAGCAATATC
51  GTTGCTGCAA ATCCCCATTA GTCATGCGAA CGGTTTGGAT GCCCGTTTGC
101 GCGATGATAT GCAGGCAAAA CACTACGAAC CGGTTGGTAA ATACCATCTG
30  151 TTTGGTAATG CTCGCGGCAG TGTAAAAAAG CGGTTTACG CCGTCCAGAC
201 ATTTGATGCA ACTGCGGTCA GTCCTGTACT GCCTATTACA CACGAACGGA
251 CAGGGTTTGA AGGTGTTATC GGTATGAAA CCCATTTTTC AGGGCACGGA
301 CATGAAGTAC ACAGTCCGTT CGATCATCAT GATTCAAAAA GCACTTCTGA
351 TTTCAGCGGC GGTGTAGACG GCGGTTTAC TGTTTACCAA CTTTCATCGAA
35  401 CAGGTCGGA AATCCATCCG GAGGATGGAT ATGACGGGCC GCAAGGCAGC
451 GATTATCCGC CCCCAGGAGG AGCAAGGGAT ATATACAGCT ATTATGTCAA
501 AGGAACCTCA ACAAACAAAG AACTAATAT TGTCCCTCAA GCCCATTTT
551 CAGACCGTTG GCTAAAGAA AATGCCGGTG CCGCTCTGG TTTTTCAGC
601 CGTGCAGATG AAGCAGGAAA ACTGATATGG GAAAGCGACC CCAATAAAAA
40  651 TTGGTGGGCT AACCGTATGG ATGATGTTCC CGGCATCGTC CAAGGTGCGG
701 TTAATCCTTT TTTAATGGGT TTTCAAGGAG TAGGATTTGG GGCAATTACA
751 GACAGTGCAG TAAGCCCGGT CACAGATACA GCCGCGCAGC AGACTCTACA
801 AGGTATTAAT GATTAGGAA AATTAAGTCC GGAAGCACAA CTTGCTGCCG
851 CGAGCCTATT ACAGGACAGT GCTTTTGGCG TAAAAGACGG TATCAACTCT
45  901 GCCAAACAAT GGGCTGATGC CCATCCAAAT ATAACAGCTA CTGCCCAAAC
951 TGCCCTTTCC GCAGCAGAGG CCGCAGGTAC GGTTTGGAGA GGTAAAAAAG
1001 TAGAACTTAA CCCGACTAAA TGGGATTGGG TTAATAATAC CGGTTATAAA
1051 AAACCTGCTG CCCGCCATAT GCAGACTTTA GATGGGGAGA TGGCAGGTGG
1101 GAATAAACCT ATTAATCTTT TACCAAACAG TGCCGCTGAA AAAAGAAAAAC
50  1151 AAAATTTTGA GAAGTTTAAT AGTAACTGGA GTTCAGCAAG TTTTGATTCA

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1	ATGAATTNGC	CTATTCAAAA	ATTTCATGATG	CTGTTTGCAG	CAGCAATATC
51	GTNGCTGCAA	ATCCCNATTA	GTCATGCGAA	CGGTTTGAT	GCCCGTTTGC
101	GCGATGATAT	GCAGGCAAAA	CAC TACGAAC	CGGGTGGTAA	ATACCATCTG
151	TTTGGTAAATG	CTCGCGCGAC	TGTTAAAAAT	CGGGTTTACG	CGCTCCAAAC
201	ATTTGATGCA	ACTGCGGTCG	GCCCCATACT	GCCTATTACA	CACGAACGGA
251	CAGGATTTTGA	AGGCATCTATC	GGTTATGAA	CCGATTTTTC	AGGACATTTGA
301	CATGAAGTAC	ACAGTCCGTT	CGATAATCAT	GATTTCAAAA	GCACCTCTGA
351	TTTCAGCGCG	GGCGTAGACG	GTGTTTTTAC	CGTTTACCAC	CTTCATCGGA
401	CAGGGTCGGA	AATCCATCCG	GAGGATGGAT	ATGACGGGCC	GCAAGGCAGC
451	GATTATCCGC	CCCCCGGAGG	AGCAAGGGAT	ATATACANNT	ANATATGTCAA
501	AGGAACCTTCA	ACAAAACAAA	AGAGTAAAT	TGTTCCCCGA	GCCCCATTTT
551	CAGACCGCTG	GCTAAAAGAA	AATGCCGGTG	CCGGCTCTGG	TTTTTTTCAGC
601	CTGTGCTGATG	AAGCAGGAAA	ACTGATATGG	GAAAGCGACC	CCAATAAAAA
651	TTGGTGGGCT	AACCGTATGG	ATGATATTCC	CGGCATCTGC	CAGGCTCGGG
701	TTAATCTCTT	TTTAATGGGT	TTTCAAGGAG	TAGGGATTGG	GCAGATTACA
751	GACAGTGCAG	TAAGCCCGGT	CACAGATACA	GCCGCGCAGC	AGACTCTACA
801	AGGTATNAAT	CATTTAGGAA	ANTTTAAGTC	CGAAGCACAA	CTTGCGGCTG
851	CACCCGCATT	ACAAGACGAT	GCTTTTGCCG	TAAAAGACGG	TATCAATTTCC
901	GCCACGCAAT	GGGCTGATGC	CCATCCGAAT	ATAACTGCAA	CAGCCCAAA

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951 TGCCCTTGCC GTAGCAGANG CCGCAACTAC GGT TTGGGGC GGTA AAAAAG
 1001 TAGAACTTAA CCCGACCAAA TGGGATTGGG TTA AAAATAC NGGCTATAAN
 1051 ACACCTGCTG TTCGCACCAT GCATACTTTG GATGGGGAAA TGGCCGGTGG
 1101 GAATAGACCG CCTAAATCTA TAACGTCCAA CAGCAAAGCA GATGCTTCCA
 1151 CACAACCGTC TTTACAAGCG CAACTAATTG GAGACAAAT TANNNNNGG
 1201 CATGCTTATA ACAAGCATGT CATAAGACAA CAAGAATTTA CGGATTTAAA
 1251 TATCAATTCA CCAGCAGATT TTGCTCGGCA TATGAAAAT ATTGTTAGCC
 1301 ATCCANCAA TATGAAAGAG TTACCTCGCG GTAGAACTGC GTATTGGGAT
 1351 NATAAAACAG GGACNATAGT TATCCGAGAT AAAAATTCTG ACGATGGAGG
 1401 TACAGCATTT AGACCAACAT CAGGTAAAAA ATATTATGAT GATTATAG

This encodes a protein having amino acid sequence <SEQ ID 168>:

1 MNXPIQKFMM LFAAAISXLQ IPISHANGLD ARLRDDMQAK HYEPGGKYHL
 51 FGNARGSVKN RVYAVQTFDA TAVGPILPIT HERTGFEGII GYETHFSGHG
 101 HEVHSPFDNH DSKSTSDFSG GVDGGFTVYQ LHRTGSEIHP EDGYDGPQGS
 151 DYPPPGGARD IYXXYVKGTS TKTKSNIVER APFSDRWLKE NAGAASGFFS
 201 RADEAGKLIW ESDPNKNWWA NRMDDIRGIV QGAVNPFLMG FQGVGIGAIT
 251 DSAVSPVTD TAAQOTLQGXN HLGXLSPEAQ LAAATALQDS AFAVKDGINS
 301 ARQWADAHPN ITATAQTALA VAXAATTWVG GKKVELNPTK WDWVKNTGYX
 351 TPAVRTMHTL DGEMAGGNRP PKSITSNSKA DASTQPSLQA QLIGE QIXXG
 401 HAYNKHVIRQ QEFTDLNINS PADFARHIEN IVSHFXNMKE LPRGRTAYWD
 451 KKTGTIVIRD KNSDDGGTAF RPTSGKKYYD DL*

ORF29a and ORF29-1 show 90.1% identity in 385 aa overlap:

		10	20	30	40	50	60
25	orf29a.pep	MNXPIQKFMM	LFAAAISXLQ	IPISHANGLD	ARLRDDMQAK	HYEPGGKYHL	FGNARGSVKN
	orf29-1	MNLPIQKFMM	LFAAAISLLQ	IPISHANGLD	ARLRDDMQAK	HYEPGGKYHL	FGNARGSVKK
		10	20	30	40	50	60
30	orf29a.pep	RVYAVQTFDA	TAVGPILPIT	HERTGFEGII	GYETHFSGHG	HEVHSPFDNH	DSKSTSDFSG
	orf29-1	RVYAVQTFDA	TAVSPVLPIT	HERTGFEGVI	GYETHFSGHG	HEVHSPFDNH	DSKSTSDFSG
		70	80	90	100	110	120
35	orf29a.pep	GVDGGFTVYQ	LHRTGSEIHP	EDGYDGPQGS	DYPPPGGARD	IYXXYVKGTS	TKTKSNIVER
	orf29-1	GVDGGFTVYQ	LHRTGSEIHP	EDGYDGPQGS	DYPPPGGARD	IYXXYVKGTS	TKTKSNIVER
		130	140	150	160	170	180
40	orf29a.pep	APFSDRWLKE	NAGAASGFFS	RADEAGKLIW	ESDPNKNWWA	NRMDDIRGIV	QGAVNPFLMG
	orf29-1	APFSDRWLKE	NAGAASGFFS	RADEAGKLIW	ESDPNKNWWA	NRMDDIRGIV	QGAVNPFLMG
		190	200	210	220	230	240
45	orf29a.pep	FQGVGIGAIT	DSAVSPVTD	TAAQOTLQGXN	HLGXLSPEAQ	LAAATALQDS	AFAVKDGINS
	orf29-1	FQGVGIGAIT	DSAVSPVTD	TAAQOTLQGXN	HLGXLSPEAQ	LAAATALQDS	AFAVKDGINS
		250	260	270	280	290	300
50	orf29a.pep	ARQWADAHPN	ITATAQTALA	VAXAATTWVG	GKKVELNPTK	WDWVKNTGYX	TPAVRTMHTL
	orf29-1	ARQWADAHPN	ITATAQTALA	VAXAATTWVG	GKKVELNPTK	WDWVKNTGYX	TPAVRTMHTL
		310	320	330	340	350	360
55	orf29a.pep	DGEMAGGNRP	PKSITSNSKA	DASTQPSLQA	QLIGE QIXXG	HAYNKHVIRQ	QEFTDLNINS
	orf29-1	DGEMAGGNRP	PKSITSNSKA	DASTQPSLQA	QLIGE QIXXG	HAYNKHVIRQ	QEFTDLNINS
		370	380	390	400	410	420
60	orf29a.pep	DGEMAGGNRP	PKSITSNSKA	DASTQPSLQA	QLIGE QIXXG	HAYNKHVIRQ	QEFTDLNINS
	orf29-1	DGEMAGGNRP	PKSITSNSKA	DASTQPSLQA	QLIGE QIXXG	HAYNKHVIRQ	QEFTDLNINS
		370	380	390	400	410	420

Homology with a predicted ORF from *N.gonorrhoeae*

ORF29 shows 88.8% identity over a 125aa overlap with a predicted ORF (ORF29.ng) from *N.*

gonorrhoeae:

5	orf29.pep	VSPVLPITHERTGFEGVIGYETHFSGHGHE	30
	orf29ng	EPGGKYHLFGNARGSVKNRVCVQTFDATAVGPILPITHERTGFEGVIGYETHFSGHGHE	102
10	orf29.pep	VHSPFDHHDHSDKSTSDFSGGVDGGFTVYQLHRTWSEIHPEDGYDGPQAAAXYPPPGGARDIY	90
	orf29ng	VHSPFDNHSDKSTSDFSGGVDGGFTVYQLHRTGSEIHPEDGYDGPQGGGYPPPGGARDIY	162
	orf29.pep	SYVVKGTSTKTKTSIVPQAPFSDRWLEENAGAASG	125
	orf29ng	SYHIKGTSTKTKINTVPQAPFSDRWLKENAGAASGFLSRADEAGKLIWENDPKNWRANR	222

- 15 The complete length ORF29ng nucleotide sequence <SEQ ID 169> is predicted to encode a protein having amino acid sequence <SEQ ID 170>:

	1	MNLPIQKFMM	LFAAAISLLQ	IPISHANGLD	ARLRDDMQAK	HYEPGGKYHL
	51	FGNARGSVKN	RVCAVQTFDA	TAVGPILPIT	HERTGFEQVI	GYETHFSGHG
20	101	HEVHSPFDNH	DSKSTSDFSG	GVDGGFTVYQ	LHRTGSEIHP	EDGYDGPQGG
	151	GYPPPGGARD	IYSYHIKGT	TKTKINTVPQ	APFSDRWLKE	NAGAAAGFLS
	201	RADEAGKLIW	ENDPDKNWRA	NRMDDIRGIV	QGAVNPFITG	FQGLGVGAIT
	251	DSAVSPVTYA	AARKTLQGIH	NLGNLSPEAQ	LAAATALQDS	AFAVKDSINS
	301	ARQWADAHNP	ITATAQTALA	VTEAATTVWG	GKKVELNPAK	WDWVKNTGYK
	351	KPAARHMQTV	DGEMAGGNKP	LESKNTVTTN	NFFENTGYTE	KVLRQASNGD
25	401	YHGFQSVSDA	FSENGTVIQI	VGGDNIVRHK	LYIPGSYKKG	DGNFEYIREA
	451	DGKINHRLFV	PNQQLPEK*			

In a second experiment, the following DNA sequence <SEQ ID 171> was identified:

	1	atgAATTTGC	CTATTCAAAA	ATTCATGATG	ctgttggcAg	cggcaatatac
	51	gatgctGCat	ATCCCCATTA	GTCATGCGAA	CGGTTGGAT	GCCCGTTTGC
30	101	GCGATGATAT	GCAGGCAAAA	CACTACGAAC	CGGGTGGCAA	ATACCATCTG
	151	TTTGGTAATG	CTCGCGGCAG	TGTTAAAAAT	CGGGTTTGGC	CCGTCCAAAC
	201	ATTTGATGCA	ACTGCGGTTCG	GCCCCATACT	GCCTATTACA	CACGAACGGA
	251	CAGGATTTGA	AGGTGTTATC	GGCTATGAAA	CCCATTTTTC	AGGACACGGA
	301	CACGAAGTAC	ACAGTCCGTT	CGATAATCAT	GATTCAAAAA	GCACCTCTGA
35	351	TTTCAGCGGC	GGCGTAGACG	GCGGTTTTC	CGTTTACCAA	CTTCATCGGA
	401	CAGGGTCGGA	AATACATCCC	GCAGACGGAT	ATGACGGGCC	TCAAGCGGCG
	451	GGTTATCCGG	AACCACAAGG	GGCAAGGGAT	ATATACAGCT	ACCATATCAA
	501	AGGAACCTCA	ACCAAAACAA	AGATAAACAC	TGTTCCGCAA	GCCCTTTTTT
	551	CAGACCGCTG	GCTAAAAAGAA	AATGCCGGTG	CCGCTTCCGG	TTTTCTCAGC
40	601	CGTGCAGATG	AAGCAGGAAA	ACTGATATGG	GAAAACGACC	CCGATAAAAA
	651	TTGGCGGGCT	AACCGTATGG	ATGATATTCG	CGGCATCGTC	CAAGGTGCGG
	701	TTAATCCTTT	TTTAACGGGT	TTTCAAGGGG	TAGGGATTGG	GGCAATTACA
	751	GACAGTGGCG	TAAGCCCGGT	CACAGATACA	GCCGCTCAGC	AGACTCTACA
	801	AGGTATTAAT	GATTTAGGAA	ATTTAAGTCC	GGAAGCACAA	CTTGCCGCGG
45	851	CGAGCCTATT	ACAGGACAGT	GCCTTTGCGG	TAAAAGACGG	CATCAATTCC
	901	GCCAGACAAT	GGGCTGATGC	CCATCCGAAT	ATAACAGCAA	CAGCCCAAAC
	951	TGCCCTTGCC	GTAGCAGAGG	CCGCAGGTAC	GGTTTGGCGC	GGTAAAAAAG
	1001	TAGAACTTAA	CCCGACCAAA	TGGGATTGGG	TTAAAAATAC	CGGCTATAAA
	1051	AAACCTGCTG	CCCGCCATAT	GCAGACTGTA	GATGGGGAGA	TGGCAGGGGG
50	1101	GAATAGACCG	CCTAAATCTA	TAACGTCGGA	AGGAAAAGCT	AATGCTGCAA
	1151	CCTATCCTAA	GTTGGTTAAT	CAGCTAAATG	AGCAAAACTT	AAATAACATT
	1201	GCGGCTCAAG	ATCCAAGATT	GAGTCTAGCT	ATTCATGAGG	GTAAAAAATA
	1251	TTTTCCAATA	GGAAGTCAA	CTTATGAAGA	GGCAGATAGA	CTAGGTAAAA
	1301	TTTGGGTGG	TGAGGGTGCA	AGACAAACTA	GTGGAGGCGG	ATGGTTAAGT
55	1351	AGAGATGGCA	CTCGACAATA	TCGGCCACCA	ACAGAAAAAA	AATCACAATT
	1401	TGCAACTACA	GGTATTCAAG	CAAAATTTGA	AACTTATACT	ATTGATTCAA
	1451	ATGAAAAAAG	AAATAAAATT	AAAAATGGAC	ATTTAAATAT	TAGGTAA

This encodes a protein having amino acid sequence <SEQ ID 172; ORF29ng-1>:

60	1	MNLPIQKFMM	LLAAAISMLH	IPISHANGLD	ARLRDDMQAK	HYEPGGKYHL
	51	FGNARGSVKN	RVCAVQTFDA	TAVGPILPIT	HERTGFEQVI	GYETHFSGHG

101 HEVHSPFDNH DSKSTSDFSG GVDGGFTVYQ LHRTGSEIHP ADGYDGPQGG
 151 GYPEPQGARD IYSYHIKGTSTKTKINTVPQ APFSDRWLKE NAGAASGFLS
 201 RADEAGKLIW ENDPKKNWRA NRMDDIRGIV QGAVNPFLTQ FQGVGIGAIT
 251 DSAVSPVTD TAAQOTLQGIN DLGNLSPEAQ LAAASLLQDS AFAVKDGINS
 301 ARQWADAHFN ITATAQTALA VAEAAGTVWR GKKVELNPTK WDWKNTGYK
 351 KPAARHMOTV DGEMAGGNRP PKSITSEGKA NAATYPKLVN QLNEQNLNNI
 401 AAQDPRLSLA IHEGKKNFPI GTATYEEADR LGKIWVGEGA RQTSGGGWLS
 451 RDGTRQYRPP TEKKSQFATT GIQANFETYT IDSNEKRNKI KNGHLNIR*

ORF29ng-1 and ORF29-1 show 86.0% identity in 401 aa overlap:

10	orf29ng-1.pep	MNLPIQKFMMLLAAAISMLHIPISHANGLDARLRDDMQAKHYEPGGKYHLFGNARGSVKN
	orf29-1	MNLPIQKFMMLFAAAISLLQIPISHANGLDARLRDDMQAKHYEPGGKYHLFGNARGSVKK
15	orf29ng-1.pep	RVCAVQTFDATAVGPILPITHERTGFEGVIGYETHFSGHGHEVHSPFDNHD SKSTSDFSG
	orf29-1	RVYAVQTFDATAVSPVLPITHERTGFEGVIGYETHFSGHGHEVHSPFDHHD SKSTSDFSG
20	orf29ng-1.pep	GVDGGFTVYQLHRTGSEIHPADGYDGPQGGGYPEPQGARDIYSYHIKGTSTKTKINTVPQ
	orf29-1	GVDGGFTVYQLHRTGSEIHPEDGYDGPQGS DYPPPGGARDIYSYVKGSTKTKTNIVPQ
25	orf29ng-1.pep	APFSDRWLKENAGAASGFLSRADEAGKLIWENDPKKNWRANRMDDIRGIVQGAVNPFLTQ
	orf29-1	APFSDRWLKENAGAASGFFSRADEAGKLIWESDPKNKNWANRMDDVRGIVQGAVNPFLMG
30	orf29ng-1.pep	FQGVGIGAITDSAVSPVTD TAAQOTLQGINDLGNLSPEAQ LAAASLLQDS AFAVKDGINS
	orf29-1	FQGVGIGAITDSAVSPVTD TAAQOTLQGINDLGKLSPEAQ LAAASLLQDS AFAVKDGINS
35	orf29ng-1.pep	ARQWADAHFNITATAQTALAVAEAAGTVWRGKKVELNPTKWDWKNTGYKKPAARHMOTV
	orf29-1	AKQWADAHFNITATAQTALSAEAGTVWRGKKVELNPTKWDWKNTGYKKPAARHMOTL
40	orf29ng-1.pep	DGEMAGGNRPKPSI-TSEGKANAAATYPKLVNQLNEQNLNNIAAQDPRLSLAIHEGKKNFP
	orf29-1	DGEMAGGNKPIKSLPNSAAEK RKQNF EKFN SNWSSAS FDSVHKTLTPNAPGILSPDKVKT
45	orf29ng-1.pep	IGTATYEEADRLGKIWVGEGARQTSGGGWLSRDGTRQYRPPTEKKSQFATTGIQANFETY
	orf29-1	RYTSLDGKITIIKDNENNYFRIHDNSRKQYLD SNGNAVKTGNLQKGKQAKDY LQQQTHIRN

Based on this analysis, including the presence of a putative leader sequence in the gonococcal protein, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 21

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 173>:

-148-

```

1  ATGAAAAAAC AAATCACCGC AGCCGTAATG ATGCTGTCTA TGATTGCCCC
51 CGCAATGGCA AACGGCTTGG ACAATCAGGC ATTTGAAGAC CAAATGTTCC
101 ACACGCGGGC AGATGCACCG ATGCAG...

```

This corresponds to the amino acid sequence <SEQ ID 174; ORF30>:

```

5      1  MKKQITAAVM MLSMIAPAMA NGLDNQAFED QMFHTRADAP MQ..

```

Further work revealed the complete nucleotide sequence <SEQ ID 175>:

```

1  ATGAAAAAAC AAATCACCGC AGCCGTAATG ATGCTGTCTA TGATTGCCCC
51 CGCAATGGCA AACGGCTTGG ACAATCAGGC ATTTGAAGAC CAAGTGTTCC
101 ACACGCGGGC AGATGCACCG ATGCAGTTGG CGGAGCTTTC TCAAAAGGAG
151 ATGAAGGAGA CAGAGGGGGC GTTCTCTCCA TTGGCTATCT TGGGTGGTGC
201 TGCCATTGGT ATGTGGACAC AGCATGGTTT TAGTTATGCA ACGACAGGCA
251 GACCAGCTTC TGTTAGAGAT GTTGCTATTG CTGGCGGATT AGGCGCAATT
301 CCTGGTGGTG TAGGCGCCGC AGGAAAGGTT GTTTCCTTTG CTAAATATGG
351 ACGTGAGATT AAAATCGGCA ATAATATGCG GATAGCCCCC TTCGGTAATA
15 401 GAACAGGTCA TCCTATTGGA AAATTCCCCC ATTATCATCG TCGAGTTACG
451 GATAATACGG GCAAGACTTT GCCTGGACAG GGAATTGGTC GTCATCGCCC
501 TTGGGAATCA AAATCTACGG ACAGATCATG GAAAAACCGC TTCTAA

```

This corresponds to the amino acid sequence <SEQ ID 176; ORF30-1>:

```

20      1  MKKQITAAVM MLSMIAPAMA NGLDNQAFED QVFHTRADAP MQLAELSQKE
51  MKETEGAFLE LAILGGAAIG MWTQHGFSA TTGRPASVRD VAIAGGLGAI
101 PGGVGAAGKV VSAKYGREI KIGNNMRIAP FGNRTGHPIG KFPYHRRVT
151 DNTGKTLPGQ GIGRHRPWES KSTDRSWKNR F*

```

Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

25 ORF30 shows 97.6% identity over a 42aa overlap with an ORF (ORF30a) from strain A of *N. meningitidis*:

```

      10      20      30      40
orf30.pep  MKKQITAAVMMLSMIAPAMANGLDNQAFEDQMFHTRADAPMQ
30 orf30a   MKKQITAAVMMLSMIAPAMANGLDNQAFEDQVFHTRADAPMQLAELSQKEMKXTXGAFLP
      10      20      30      40      50      60
orf30a     LXILGGAAIGMWTQHGFSAATTGRPASVRDVAIAGGLGAIPGXVGAAGKVVSFAKYGREI
      70      80      90     100     110     120

```

35 The complete length ORF30a nucleotide sequence <SEQ ID 177> is:

```

1  ATGAAAAAAC AAATCACCGC AGCCGTAATG ATGCTGTCTA TGATTGCCCC
51 CGCAATGGCA AACGGCTTGG ACAATCAGGC ATTTGAAGAC CAAGTGTTCC
101 ACACGCGGGC AGATGCACCG ATGCAGTTGG CGGAGCTTTC TCAAAAGGAG
151 ATGAAGGANA CAGNGGGGGC GTTCTCTCCA TTGGNTATCT TGGGTGGTGC
40 201 TGCCATTGGT ATGTGGACAC AGCATGGTTT TAGTTATGCA ACGACAGGCA
251 GACCAGCTTC TGTTAGAGAT GTTGCTATTG CTGGCGGATT AGGCGCAATT
301 CCTGGTGNTG TAGGCGCCGC AGGAAAGGTT GTTTCCTTTG CTAAATATGG
351 ACGTGAGATT AAAATCGGCA ATAATATGCG GATAGCCCCC TTCGGTAATA
401 GAACAGGTCA TCCTATTGGA AAATTCCCCC ATTATCATCG TCGAGTTACG
45 451 GATAATACGG GCAAGACTTT GCCTGGACAG GGAATTGGTC GTCATCGCCC
501 TTGGGAATCA AAATCTACGG ACAGATCATG GAAAAACCGC TTCTAA

```

This encodes a protein having amino acid sequence <SEQ ID 178>:

```

50      1  MKKQITAAVM MLSMIAPAMA NGLDNQAFED QVFHTRADAP MQLAELSQKE
51  MKXTXGAFLP LXILGGAAIG MWTQHGFSA TTGRPASVRD VAIAGGLGAI
101 PGXVGAAGKV VSAKYGREI KIGNNMRIAP FGNRTGHPIG KFPYHRRVT
151 DNTGKTLPGQ GIGRHRPWES KSTDRSWKNR F*

```

ORF30a and ORF30-1 show 97.8% identity in 181 aa overlap:

```

orf30a.pep  MKKQITAAVMMLSMIAPAMANGLDNQAFEDQVFHTRADAPMQLAELSQKEMKXTXGAFLP 60

```

```

      ||||||||||||||||||||||||||||||||||||||||||||||||||||
orf30-1  MKKQITAAVMMLSMIAPAMANGLDNQAFEDQVFHTRADAPMQLAELSQKEMKETEGAFLP  60
5  orf30a.pep  LXILGGAAGMWTQHGFYSYATTGRPASVRDVAIAGGLGAIPGXVGAAGKVVSFAKYGREI 120
      ||||||||||||||||||||||||||||||||||||||||||||||||||||
orf30-1  LAILGGAAGMWTQHGFYSYATTGRPASVRDVAIAGGLGAIPGGVGAAGKVVSFAKYGREI 120
      ||||||||||||||||||||||||||||||||||||||||||||||||||||
orf30a.pep  KIGNNMRIAPFGNRTGHPICKFPHYHRRVTDNTGKTLPGQIGRHRPWESKSTDRSWKNR 180
10 orf30-1  KIGNNMRIAPFGNRTGHPICKFPHYHRRVTDNTGKTLPGQIGRHRPWESKSTDRSWKNR 180
      ||||||||||||||||||||||||||||||||||||||||||||||||||||
orf30a.pep  FX
      ||
orf30-1  FX
15

```

Homology with a predicted ORF from *N.gonorrhoeae*

ORF30 shows 97.6% identity over a 42aa overlap with a predicted ORF (ORF30.ng) from *N. gonorrhoeae*:

```

20 orf30.pep  MKKQITAAVMMLSMIAPAMANGLDNQAFEDQMFHTRADAPMQ 42
      ||||||||||||||||||||||||||||||||||||||||||||||||||||
orf30ng  MKKQITAAVMMLSMIAPAMANGLDNQAFEDQVFHTRADAPMQLAELSQKEMKETEGAFLP  60

```

The complete length ORF30ng nucleotide sequence <SEQ ID 179> is

```

      1  ATGAAAAAAC  AAATCACCGC  AGCCGTAATG  ATGCTGTCTA  TGATCGCCCC
25  51  CGCAATGGCA  AACGGATTGG  ACAATCAGGC  ATTGAAGAC  CAAGTGTGCC
      101  ACACGCGGGC  AGATGCGCCG  ATGCAGTTGG  CGGAGCTTTC  TCAGAAGGAG
      151  ATGAAGGAGA  CTGAAGGGGC  TTTCTTCCA  TTGGCTATCT  TGGGTGGTGC
      201  TGCCATTGGT  ATGTGGACAC  AGCATGGTTT  TAGTTATGCA  ACGACAGGCA
      251  GACCAGCTTC  TGTTAGAGAT  GTTGCTGGCG  GATTAGGCGC  AATTCCTGGT
      301  GATGTAGGTG  CTGCAGGAAA  GGTGTTTCC  TTTGCTAAAT  ATGGACGTGA
30  351  GATTAAAAATC  GGCAATAATA  TCGGATAGC  CCCTTTCGGT  AATAGAACAG
      401  GTCATCCTAT  TGGAAAATTT  CCCCATTATC  ATCGTCGAGT  TACGGATAAT
      451  ACGGGCAAGA  CTTGCGCTGG  ACAGGGAATT  GGTCGTCATC  GCCCTTGGGA
      501  ATCAAAATCT  ACGGACAGAT  CATGGAAAAA  CCGCTTCTAA

```

This encodes a protein having amino acid sequence <SEQ ID 180>:

```

35  1  MKKQITAAVM  MLSMIAPAMA  NGLDNQAFED  QVFHTRADAP  MQLAELSQKE
      51  MKETEGAFLP  LAILGGAAG  MWTQHGFSA  TTGRPASVRD  VAGGLGAIPG
      101  DVGAAGKVVS  FAKYGREIKI  GNNMRIAPFG  NRTGHPICKF  PHYHRRVTDN
      151  TGKTLPGQGI  GRHRPWESKS  TDRSWKNRF*

```

ORF30ng and ORF30-1 show 98.3% identity in 181 aa overlap:

```

40      10      20      30      40      50      60
orf30ng.pep  MKKQITAAVMMLSMIAPAMANGLDNQAFEDQVFHTRADAPMQLAELSQKEMKETEGAFLP
      ||||||||||||||||||||||||||||||||||||||||||||||||||||
orf30-1  MKKQITAAVMMLSMIAPAMANGLDNQAFEDQVFHTRADAPMQLAELSQKEMKETEGAFLP
      10      20      30      40      50      60
45      70      80      90      100     110
orf30ng.pep  LAILGGAAGMWTQHGFYSYATTGRPASVRDVA--GGLGAIPGDVGAAGKVVSFAKYGREI
      ||||||||||||||||||||||||||||||||||||||||||||||||||||
orf30-1  LAILGGAAGMWTQHGFYSYATTGRPASVRDVAIAGGLGAIPGGVGAAGKVVSFAKYGREI
50      70      80      90      100     110     120
      120     130     140     150     160     170
orf30ng.pep  KIGNNMRIAPFGNRTGHPICKFPHYHRRVTDNTGKTLPGQIGRHRPWESKSTDRSWKNR
      ||||||||||||||||||||||||||||||||||||||||||||||||||||
55 orf30-1  KIGNNMRIAPFGNRTGHPICKFPHYHRRVTDNTGKTLPGQIGRHRPWESKSTDRSWKNR
      130     140     150     160     170     180
      180
orf30ng.pep  FX
      ||
60 orf30-1  FX

```

Based on this analysis, including the presence of a putative leader sequence in the gonococcal protein, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 22

- 5 The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 181>:

```

1 ATGAATAAAA CTCTCTATCG TGTAATTTTC AACCGCAAAC GTGGGGCTGT
51 GrTAGCCGTT GCTGAAACTA CCAAGCGCGA AGGTAAAAGC TGTGCCGATA
101 GTGATTCAGG CAGCGCTCAT GTGAAATCTG TTCCTTTTGG TACTACTCAT
151 GCACCTGTTT GTg.CGTtAc AAATATCTTT TCTTTTCTT TATTGGGCTT
10 201 TTCTTTATGT TTGGCTGTAG GtacGGyCA TATTGCTTTT GCTGATGGCA
251 TT..

```

This corresponds to the amino acid sequence <SEQ ID 182; ORF31>:

```

1 MNKTLRYVIF NRKRGAVXAV AETTKREGKS CADSDSGSAH VKSVFPFGTTH
51 APVCXVTNIF SFSLLGFSLC LAVGTXNIAF ADGI..

```

- 15 Further work revealed a further partial nucleotide sequence <SEQ ID 183>:

```

1 ATGAATAAAA CTCTCTATCG TGTAATTTTC AACCGCAAAC GTGGGGCTGT
51 GGTAGCCGTT GCTGAAACTA CCAAGCGCGA AGGTAAAAGC TGTGCCGATA
101 GTGATTCAGG CAGCGCTCAT GTGAAATCTG TTCCTTTTGG TACTACTCAT
151 GCACCTGTTT GTCGTTCAAA TATCTTTTCT TTTTCTTAT TGGGCTTTTC
20 201 TTTATGTTTG GCTGTAGGTA CGGCCAATAT TGCTTTTGCT GATGGCATT..

```

This corresponds to the amino acid sequence <SEQ ID 184; ORF31-1>:

```

1 MNKTLRYVIF NRKRGAVVAV AETTKREGKS CADSDSGSAH VKSVFPFGTTH
51 APVCRSNIFS FSLGLFSLCL AVGTANIAFA DGI..

```

Computer analysis of this amino acid sequence gave the following results:

- 25 Homology with a predicted ORF from *N.gonorrhoeae*

ORF31 shows 76.2% identity over a 84aa overlap with a predicted ORF (ORF31.ng) from *N.gonorrhoeae*:

```

30 orf31.pep      MNKTLRYVIFNRKRGAVXAVAETTKREGKSCADSDSGSAHVKSVPFGTTHAPVCXVTNIF 60
   |||||
orf31ng         MNKTLRYVIFNRKRGAVVAVAVETTKREGKSCADSGSGSVYVKSVSFIPTH-----SKAF 54

orf31.pep      SFSLLGFSLC LAVGTXNIAFADGI 84
   || |||||
orf31ng         CFSALGFSLC LALGTVNIAFADGIITDKAAPKTQQATILQTGNGIPQVNIQTPTSAGVSV 114

```

- 35 The complete length ORF31ng nucleotide sequence <SEQ ID 185> is:

```

1 ATGAACAAAA CCCTCTATCG TGTGATTTTC AACCGCAAAC GCGGTGCTGT
51 GGTAGCTGTT GCCGAAACCA CCAAGCGCGA AGGTAAAAGC TGTGCCGATA
101 GTGGTTCGGG CAGCGTTTAT GTGAAATCCG TTCCTTTTCAT TCCTACTCAT
151 TCCAAAGCCT TTTGTTTTTC TGCAATTAGG CTTTCTTTAT GTTTGGCTTT
40 201 GGGTACGGTC AATATTGCTT TTGCTGACGG CATTATTACT GATAAAGCTG
251 CTCCTAAAAC CCAACAAGCC ACGATTCTGC AAACAGGTaa cGGCATACCG
301 CAAGTCAATA TTCAAACCCC TACTTCGGCA GGGGTTTCTG TTAATCAATA
351 TGCCAGTTT GATGTGGGTA ATCGCGGGGC GATTTTAAAC AACAGTCGCA
401 GCAACACCCA AACACAGCTA GCGGTTGGA TTCAAGGCAA TCCTTGTTG
45 451 ACAAGGGGCG AAGCACGTGT GGTGTAAAC CAAATCAACA GCAGCCATCC
501 TTCACAACTG AATGGCTATA TTGAAGTGGG TGGACGACGT GCAGAAGTCG
551 TTATTGCCAA TCCGGCAGGG ATGTCAGTCA ATGGTGGTGG TTTTATCAAT
601 GCTTCCCGTG CCACTTTGAC GACAGGCCAA CCGCAATATC AAGCAGGAGA
651 CTTTAGCGGC TTTAAGATAA GGCAAGGCAA TGCTGTAATC GCCGGACACG

```

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701 GTTGGATGC CCGTGATACC GATTTCACAC GTATTCTTGT ATGCCAACAA
 751 AATCACCTTG ATCAGTACGG CCGAACAAGC AGGCATTTCGT AA

This encodes a protein having amino acid sequence <SEQ ID 186>:

5 1 MNKTLYRVIF NRKRGAVVAV AETTKREGKS CADSGSGSVY VKSVSFIPTH
 51 SKAFCFSALG FSLCLALGTV NIAFADGIIT DKAAPKTQQA TILQTGNGIP
 101 QVNIQTPTSA GVSVNQYAQF DVGNRGAILN NSRSNTQTQL GGWIQGNPWL
 151 TRGEARVVVN QINSSHPSQL NGYIEVGGRR AEVVIANPAG IAVNGGGFIN
 201 ASRATLTGQ PQYQAGDFSG FKIRQGNNAVI AGHGLDARDT DFTRILVCQQ
 251 NHLDQYGRTS RHS*

10 This gonococcal protein shares 50% identity over a 149aa overlap with the pore-forming hemolysins-like HecA protein from *Erwinia chrysanthemi* (accession number L39897):

orf31ng 96 GNGIPQVNIQTPTSAGVSVNQYAQFDVGNRGAILNNSRSN-TQTQLGGWIQGNPWLTRGE 154
 GNG+P VNI TP ++G+S N+Y F+V NRG ILNN + T +QLGG IQ NP L
 HecA 45 GNGVPVNIATPDASGLSHNRYHDFVNDNRGLILNNGTARLTPSQLGGLIQNNPNLNGRA 104
 15 Orf31ng 155 ARVVVNQINSSHPSQLNGYIEVGGRRAEVVIANPAGIAVNGGGFINASRATLTGQPQYQ 214
 A ++N++ S + S+L GY+EV G+ A VV+ANP GI +G GF+N R TLTTG PQ+
 HecA 105 AAAILNEVVSPNRSRLAGYLEVAGQAANVVVANPYGITCSGCGFLNTPRLTLTTGTPQFD 164
 20 Orf31ng 215 -AGDFSGFKIRQGNNAVIAGHGLDARDTDF 242
 AG SG +R G+ +I G GLDA +D+
 HecA 165 AAGGLSGLDVRGGDILIDGAGLDASRSDY 193

Furthermore, ORF31ng and ORF31-1 show 79.5% identity in 83 aa overlap:

25 orf31-1.pep 10 20 30 40 50 60
 MNKTLYRVIFNRKRGAVVAVAEETTKREGKSCADSDSGSAHVKSVPFGTTHAPVCRSNIFS
 orf31ng MNKTLYRVIFNRKRGAVVAVAEETTKREGKSCADSGSGSVYVKSVSFIPTH-----SKAFC
 10 20 30 40 50
 30 orf31-1.pep 70 80
 FSLLGFSCLAVGTANIAFADGI
 orf31ng FSALGFSCLALGTVNIAFADGIITDKAAPKTQQATILQTGNGIPQVNIQTPTSAGVSVN
 60 70 80 90 100 110

35 On this basis, including the homology with hemolysins, and also with adhesins, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 23

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 187>:

40 1 ATGAATACTC CTCCTTTTGT CTGTTGGATT TTTTGCAAGG TCATCGACAA
 51 TTTCGGCGAC ATCGGCGTTT CGTGGCGGCT CGCCCGTGT TGCACCGCG
 101 AACTCGGTTG GCAGGTGCAT TTGTGGACGG ACGATGTGTC CGCCTTGCGT
 151 GCGCTTTGCC CTGATTGCC CGATGTTCCC TGCCTTCATC AGGATATTCA
 201 TGTCCGCACT TGGCATTCCG ATGCGGCAGA TATTGATACC GCG..

45 This corresponds to the amino acid sequence <SEQ ID 188; ORF32>:

1 MNTPPFVCWI FCKVIDNFGD IGVSRLARV LHRELGWQVH LWTDDVSALR
 51 ALCPDLPDVP CVHQDIHVRT WHSDAADIDT A..

Further work revealed the complete nucleotide sequence <SEQ ID 189>:

50 1 ATGAATACTC CTCCTTTTGT CTGTTGGATT TTTTGCAAGG TCATCGACAA
 51 TTTCGGCGAC ATCGGCGTTT CGTGGCGGCT CGCCCGTGT TGCACCGCG
 101 AACTCGGTTG GCAGGTGCAT TTGTGGACGG ACGATGTGTC CGCCTTGCGT

```

151 GCGCTTTGCC CTGATTTGCC CGATGTTCCC TGC GTTCATC AGGATATTCA
201 TGTCCGCACT TGGCATTCCG ATGCGGCAGA TATTGATACC GCGCCTGTTC
251 CCGATGTCTG CATCGAAACT TTTGCCTGCG ACCTGCCCGA AAATGTGCTG
301 CACATTATCC GCCGACACAA GCCGCTTTGG CTGAATTGGG AATATTTGAG
351 CGCGGAGGAA AGCAATGAAA GGCTGCATCT GATGCCTTCG CCGCAGGAGG
401 GTGTTCAAAA ATATTTTGGG TTTATGGGTT TCAGCGAAAA AAGCGGCGGG
451 TTGATACGCG AACGTGATTA CTGCGAAGCC GTCCGTTTCG ATACTGAAGC
501 CCTGCGAGAG CGGCTGATGC TGCCCGAAAA AAACGCCTCC GAATGGCTGC
551 TTTTCGGCTA TCGGAGCGAT GTTTGGGCAA AGTGGCTGGA AATGTGGCGA
601 CAGGCAGGCA GCCCGATGAC ACTGTTGCTG GCGGGGACGC AAATCATCGA
651 CAGCCTCAA CAAAGCGGCG TTATTCGCA AGATGCCCTG CAAAACGACG
701 GCGATGTTTT TCAGACGGCA TCCGTCCGCC TCGTCAAAAT CCCTTTCGTG
751 CCGCAACAGG ACTTCGACCA ACTGCTGCAC CTTGCCGACT GCGCCGTCAT
801 CCGCGGCGAA GACAGTTTCG TGCGCGCCCA GCTTGCGGGC AAACCTTCTT
851 TTTGGCACAT CTACCCGCAA GACGAGAATG TCCATCTCGA CAAACTCCAC
901 GCCTTTTGGG ATAAGGCACA CGGTTTCTAC ACGCCCGAAA CCGTGTCCGG
951 ACACGCGCGT CTTTCGGACG ACCTCAACGG CGGAGAGGCT TTATCCGCAA
1001 CACAACGCCT CGAATGTTGG CAAACCCTGC AACAACATCA AAACGGCTGG
1051 CGGCAAGGCG CGGAGGATTG GAGCCGTTAT CTTTTCGGGC AGCCGTCAGC
1101 TCCTGAAAAA CTCGCTGCCT TTGTTTCAA GCATCAAAAA ATACGCTAG

```

This corresponds to the amino acid sequence <SEQ ID 190; ORF32-1>:

```

1 MNTPPFVCWI FCKVIDNFGD IGVSRLARV LHRELGWQVH LWTDDVSALR
51 ALCPDLPDVP CVHQDIHVRT WHSDAADIDT APVPDVVIET FACDLPENVL
101 HIIRRHKPLW LNWEYLSAEE SNERLHLMPS PQEGVQKYFW FMGFSEKSGG
151 LIRERDYCEA VRFDEALRE RLMLPEKNAS EWLLFGYRSD VWAKWLEMWR
201 QAGSPMTLLL AGTQIIDLK QSGVIPQDAL QNDGDVFTQA SVRLVKIPFV
251 PQQDFDQLLH LADCAVIRGE DSFVRAQLAG KPFFWHIYPQ DENVHLDKLH
301 AFWDKAHGFY TPETVSAHRR LSDDLNGGEA LSATQRLECW QTLQQHQNGW
351 RQGAEDWSRY LFGQPSAPEK LAAFVSKHKQ IR*w

```

Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF32 shows 93.8% identity over a 81aa overlap with an ORF (ORF32a) from strain A of *N. meningitidis*:

```

35      10      20      30      40      50      60
orf32.pep MNTPPFVCWIFCKVIDNFGDIGVSWRLARVLHRELGWQVHLWTDDVSALRALCPDLPDVP
      |||||
orf32a    MNTPPFSAGXFCKVIDNFGDIGVSWRLARVLHRELGWQVHLWTDDVSALRALCPDLPDVX
      10      20      30      40      50      60

40      70      80
orf32.pep CVHQDIHVRTWHSDAADIDTA
      |||||
orf32a    CVHQDIHVRTWHSDAADIDTAPVXDVVIETFACDLPENVLHIIRRHKPLWLXWEYLSAEX
      70      80      90      100     110     120

```

The complete length ORF32a nucleotide sequence <SEQ ID 191> is:

```

1 ATGAATACTC CTCCTTTTTC TGCTGGANTT TTTTGAAGG TCATCGACAA
51 TTTTCGGCGAC ATCGGCGTTT CGTGGCGGCT TGCCCGTGTT TTGCACCGCG
101 AACTCGGTTG GCAGGTGCAT TTGTGGACGG ACGATGTGTC CGCCTTGCGT
151 GCGCTTTGCC CTGATTTGCC CGATGTTTNC TGC GTTCATC AGGATATTCA
201 TGTCCGCACT TGGCATTCCG ATGCGGCAGA TATTGATACC GCGCCTGTTC
251 NGATGTCTG CATCGAAACT TTTGCCTGCG ACCTGCCCGA AAATGTGCTG
301 CACATCATCC GCCGACACAA GCCGCTTTGG CTGAANTGGG AATATTTGAG
351 CGCGGAGGAN AGCAATGAAA GGCTGCACNT GATGCCTTCG CCGCAGGAGA
401 GTGTTCAAAA ATANTTTTGG TTTATGGGTT TCAGCGAANN NAGCGGCGGA
451 CTGATACGCG AACGCGATTA CTGCGAAGCC GTCCGTTTCG ATAGCGGAGC
501 CTTGCGCAAG AGGCTGATGC TTCCCGAAAA AAACGNCCCC GAATGGCTGC
551 TTTTCGGCTA TCGGAGCGAT GTTTGGGCAA AGTGGCTGGA AATGTGGCGA
601 CAGGCAGGCA GTCCGTTGAC ACTTTGCTG GCNCGGCGC ANATTATCGA
651 CAGCCTCAA CAAAACGGCG TTATTCGCA AGATGCCCTG CAAAACGACG
701 GCGATGTTTT TCAGACGGCA TCCGTCCGCC TCGTCAAAAT CCCTTTCGTG
751 CCGCAACAGG ACTTCGACAA ACTGCTGCAC CTTGCCGACT GCGCCGTCAT

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5
 801 CCGCGGCGAA GACAGTTTCG TGCGGCGCCA GCTTGCGGGC AAACCCCTTCT
 851 TTTGGCACAT CTACCCGCAA GATGAGAATG TCCATCTCGA CAAACTCCAC
 901 GCCTTTTGGG ATAAGGCACA CGGTTTCTAC ACGCCCGAAA CCGCATCGGC
 951 ACACCGCCGC CTTTCAGACG ACCTCAACGG CGGAGAGGCT TTATCCGCAA
 1001 CACAACGCCT CGAATGTTGG CAAATCCTGC AACACATCA AAACGGCTGG
 1051 CGGCAAGGCG CGGAGGATTG GAGCCGTTAT CTTTGTGGG AGCCTTCCGC
 1101 ATCCGAAAAA CTCGCCGCCT TTGTTTCAA GCATCAAAAA ATACGCTAG

This encodes a protein having amino acid sequence <SEQ ID 192>:

10
 1 MNTPPFSAGX FCKVIDNFGD IGVSRLARV LHRELGWQVH LWTDDVSALR
 51 ALCPDLPDVX CVHQDIHVRT WHSDAADIDT APVXDVIET FACDLPENVL
 101 HIIIRHKPLW LXWEYLSAEX SNERLHXMP S QESVXKXFW FMGFSEXSGG
 151 LIRERDYCEA VRFDSGALRK RLMLPEKNXP EWLFGYRSD VWAKWLEMWR
 201 QAGSPLTLLL AGAXIIDSLK QNGVIPQDAL QNDGDVFQTA SVRLVKIPFV
 251 PQQDFDKLLH LADCAVIRGE DSFVRAQLAG KPFFWHIYPQ DENVHLDKLH
 301 AFWDKAHGFY TPETASAHRR LSDDLNGGEA LSATQRLECW QILQQHQNGW
 351 RQGAEDWSRY LFGQPSASEK LAAFVSKHQK IR*

ORF32a and ORF32-1 show 93.2% identity in 382 aa overlap:

20
 orf32-1.pep 10 20 30 40 50 60
 orf32a 10 20 30 40 50 60
 25
 orf32-1.pep 70 80 90 100 110 120
 orf32a 70 80 90 100 110 120
 30
 orf32-1.pep 130 140 150 160 170 180
 orf32a 130 140 150 160 170 180
 35
 orf32-1.pep 190 200 210 220 230 240
 orf32a 190 200 210 220 230 240
 40
 orf32-1.pep 250 260 270 280 290 300
 orf32a 250 260 270 280 290 300
 45
 orf32-1.pep 310 320 330 340 350 360
 orf32a 310 320 330 340 350 360
 50
 orf32-1.pep 370 380
 orf32a 370 380

60 Homology with a predicted ORF from *N.gonorrhoeae*

ORF32 shows 95.1% identity over a 82aa overlap with a predicted ORF (ORF32.ng) from *N. gonorrhoeae*:

orf32.pep MNTPPF-VCWIFCKVIDNFGDIGVSWRLARVLHRELGWQVHLWTDDVSALRALCPDLP 57
 ||| | |||||

```

      orf32ng      MVMNTYAFFVCWIFCKVIDNFGDIGVSWRLARVLHRELGWQVHLWTTDDVSALRALCPDLP      60
      orf32.pep    DVPCVHQDIHVRTWHSDAADIDTA                                          81
      ||| |||||
5      orf32ng      DVPCVHQDIHVRTWHSDAADIDTAPVDAVIETTFACDLPENVLNIIRRHKPLWLNWEYLS      120

```

An ORF32ng nucleotide sequence <SEQ ID 193> was predicted to encode a protein having amino acid sequence <SEQ ID 194>:

```

      1  MVMNTYAFFV CWIFCKVIDN FGDIGVSWRL ARVLHRELGW QVHLWTTDDVS
      51  ALRALCPDLP DVPCVHQDIH VRTWHSDAAD IDTAPVDAVI IETTFACDLP
101  NVLNIIRRHK PLWLNWEYLS AESNERLHL MPSPQEGVQK YWFWMGFSEK
151  SGGIIRERDY REAVRFDTEA LRRRLVLP EK NAPEWLLFGY RGDVWAKWLD
201  MWQQAGSLMT LLAGAQIID SLKQSGVIPQ NALQNEGGVF QTASVRLVKI
251  PFVPQQDFDK LLHLADCAVI RGEDSFVRTQ LAGKPFFWHI YPQDENVHLD
301  KLHAFWDKAY GFYTPETASV HRLSDDLNG GEALSATQRL ECGVL*

```

15 Further sequencing revealed the following DNA sequence <SEQ ID 195>:

```

      1  ATGAATACAT ACGCTTTTCC TGTCTGTTGG ATTTTTTTCGA AGGTCATCGA
      51  CAATTTTCGGC GACATCGGCG TTTCGTGGCG GCTCGCCCGT GTTTTGCACC
101  GCGAAGTCGG TTGGCAGGTG CATTTGTGGA CGGACGACGT GTCCGCCTTG
151  CGCGCGCTTT GTCCCGATTT GCCCGATGTT CCCTTCGTTC ATCAGGATAT
201  TCATGTCCGC ACTTGGCATT CCGATGCGGC AGACATTGAT ACCGCGCCCG
251  TTCCCGATGC CGTTATCGAA ACTTTGCCT GCGACCTGCC CGAAAATGTG
301  CTGAACATCA TCCGCCGACA CAAACCGCTT TGGCTGAATT GGGAATATTT
351  GAGCGCGGAG GAAAGCAATG AAAGGCTGCA CCTGATGCCT TCGCCGCGAG
401  AGGGCGTTCA AAAATATTTT TGGTTTATGG GTTTCAGCGA AAAAAGCGGC
25  451  GGGTTGATAC GCGAACGCGA TTACCGCGAA GCCGTCCGTT TCGATACCGA
501  AGCCCTGCGC CGGCGGCTGG TGCTGCCCGA AAAAAACGCC CCCGAATGGC
551  TGCTTTTCGG CTATCGGGGC GATGTTTGGG CAAAGTGGCT GGACATGTGG
601  CAACAGGCAG GCAGCCTGAT GACCCACTG CTGGCGGGGG CGCAAATTAT
651  CGACAGCCTC AAACAAAGCG GCGTTATTC GCAAAACGCC CTGCAAAAtg
30  701  aaggcgGTGT CTTTCagacG gcatcgTcC gccttGTCAA AAtcCCGTTC
751  GTGCcGCAAC AGGAcTTCGA CAAATTGCTG CAcctcgCG ACTGCGCCGT
801  GATACGCGGC GAAGACAGTT TCGTGCGTAC CCAGCTTGCC GGAAAACCTT
851  TTTTGTGGCA CATCTACCCG CAAGACGAGA ATGTCCATCT CGACAAACTC
901  CAGCCTTTT GGGATAAGGC ATACGGCTTC TACACGCCCG AAACCGCATC
35  951  GGTGCACCGC CTCCCTTCGG ACGACCTCAA CGGCGGAGAG GCTTTATCCG
1001  CAACACAACG CCTCGAATGT TGGCAAACCC TGCAACAACA TCAAAACGGC
1051  TGGCGGCAAG GCGCGGAGGA TTGGAGCCGT TATCTTTTCG GGCAGCCTTC
1101  CGATCCGAA AACTCGCGC CTTTGTTC AAAGCATCAA AAAATACGCT
1151  AG

```

40 This encodes a protein having amino acid sequence <SEQ ID 196; ORF32ng-1>:

```

      1  MNTYAFFVCW IFCKVIDNFG DIGVSWRLAR VLHRELGWQV HLWTTDDVSAL
      51  RALCPDLPDV PFVHQDIHVR TWHSDAADID TAPVDAVIE TFACDLPENV
101  LNIIRRHKPL WLNWEYLSAE ESNERLHLMP SPQEGVQKYF WFMGFSEKSG
151  GLIRERDYRE AVRFDEALR RRLVLP EKNA PEWLLFGYRG DVWAKWLDMW
45  201  QQAGSLMTLL LAGAQIIDSL KQSGVIPQNA LQNEGGVFQT ASVRLVKIPF
251  VPQQDFDKLL HLADCAVIRG EDSFVRTQLA GKPFFWHIYP QDENVHLDKL
301  HAFWDKAYGF YTPETASVHR LLSDDLNGGE ALSATQRL EC WQTLQQHQNG
351  WRQGAEDWSR YLFGQPSASE KLAAFVSKHQ KIR*

```

ORF32ng-1 and ORF32-1 show 93.5% identity in 383 aa overlap:

```

50      10      20      30      40      50      59
      orf32-1.pep  MNTPPF-VCWIFCKVIDNFGDIGVSWRLARVLHRELGWQVHLWTTDDVSALRALCPDLPDV
      ||| | |||||
      orf32ng-1   MNTYAFFVCWIFCKVIDNFGDIGVSWRLARVLHRELGWQVHLWTTDDVSALRALCPDLPDV
      10      20      30      40      50      60
55      60      70      80      90      100     110     119
      orf32-1.pep  PCVHQDIHVRTWHSDAADIDTAPVDPVVIETTFACDLPENVLHIIRRHKPLWLNWEYLSAE
      | |||||
      orf32ng-1   PFVHQDIHVRTWHSDAADIDTAPVDAVIETTFACDLPENVLNIIRRHKPLWLNWEYLSAE
60      70      80      90      100     110     120
      120     130     140     150     160     170     179

```


5	orf32-1.pep	ESNERLHLMPSPOEGVQKYFWFMGFSEKSGGLIRERDYCEAVRFDTEALRRLMLPEKNA
	orf32ng-1	ESNERLHLMPSPOEGVQKYFWFMGFSEKSGGLIRERDYREAVRFDTEALRRRLVLPEKNA
10		130 140 150 160 170 180
15	orf32-1.pep	180 190 200 210 220 230 239
	orf32ng-1	SEWLLFGYRSDVWAKWLEMRQAGSEPMTLLLAGTQIIDS LKQSGVIPQDALQNDGDVFQT
20		190 200 210 220 230 240
25	orf32-1.pep	240 250 260 270 280 290 299
	orf32ng-1	ASVRLVKIPFVPQQDFDQLHLADCAVIRGEDSFVRAQLAGKPF FWHIYPQDENVHLDKL
30		250 260 270 280 290 300
35	orf32-1.pep	300 310 320 330 340 350 359
	orf32ng-1	HAFWDKAHGFYTPETVSAHRRLSDDLNGGEALSATQRLECWQTLQQHQNGWRQGAEDWSR
40		310 320 330 340 350 360
45	orf32-1.pep	360 370 380
	orf32ng-1	YLFQGPSAPEKLA AFVSKHQKIRX
50		370 380

On this basis, including the RGD sequence in the gonococcal protein, characteristic of adhesins, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

ORF32-1 (42kDa) was cloned in pET and pGex vectors and expressed in *E.coli*, as described above. The products of protein expression and purification were analyzed by SDS-PAGE. Figure 7A shows the results of affinity purification of the His-fusion protein, and Figure 7B shows the results of expression of the GST-fusion in *E.coli*. Purified His-fusion protein was used to immunise mice, whose sera were used for ELISA, giving a positive result. These experiments confirm that ORF32-1 is a surface-exposed protein, and that it is a useful immunogen.

Example 24

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 197>:

1	..TTGTTCTCTGC	GTGTNAAAGT	GGGGCGTTTT	TTCAGCAGTC	CGGCGACGTG
51	GTTTCGGGNC	AAAGACCCTG	TAAATCAGGC	GGTGTTCGG	CTGTATNCCG
101	ACGAGTGGCG	GCA.ACTTCG	GTACGTTGGA	AAATAGNCGC	AACGTGCGAC
151	AGCCTGTGGC	TCTGCAGCT	GCTCGGAATG	CTGGTGTTCG	TATTGTTGCT
201	GCTTTTGGTG	CGGCAATATA	CGTTCAACTG	GGAAAGCACG	CTGTTGAGCA
251	ATGCCGCTTC	GGTACGCGCG	GTGAAATGT	TGGCATGGCT	GCCGTCGAAA
301	CTCGGTTTCC	CTGTCCCGA	TGCGCGGTTCG	GTCATCGAAG	GCCGTCGAA
351	CGGCAATATT	GCCGATGCGC	GGGCTTGGTC	GGGGCTGCTG	GTCGNCAGTA
401	TGCCTGCTA	NGGCATCCTG	CCGCGCCTG..		

This corresponds to the amino acid sequence <SEQ ID 198; ORF33>:

1	..LFLRVKVRG	FSSPATWFRX	KDPVNQAVLR	LYXDEWRXTS	VRWKIXATSH
51	SLWLCTLLGM	LVSLLLLLV	ROYTFNWEST	LLSNAASVRA	VEMLAWLPSK
101	LGFPVPDARS	VIEGRLNGNI	ADARAWSGLL	VXSIACXGIL	PRL..

Further work revealed the complete nucleotide sequence <SEQ ID 199>:

```

      1  ATGTTGAATC CATCCCCGAAA ACTGGTTGAG CTGGTCCGTA TTTTGGACGA
    51  AGGCGGTTTT ATTTTCAGCG GCGATCCCGT ACAGGCGACG GAGGCTTTGC
   101  GCCGCGTGGA CGGCAGTACG GAGGAAAAAA TCATCCGTCG GGCGGAGATG
  151  ATTGACAGGA ACCGTATGCT GCGGGAGACG TTGGAACGTG TGCCTGCGGG
  201  GTCGTTCTGG TTGTGGGTGG TGGCGGCGAC GTTTGCATTT TTTACCGTT
  251  TTTTCTGAC TTATCTTCTA ATGGACAATC AGGGTCTGAA TTTCTTTTGT
  301  GTTTTGGCGG GCGTGTTGGG CATGAATACG CTGATGCTGG CAGTATGGTT
  351  GGCATATGTT TTCCTGCGTG TGAAAGTGGG GCGTTTTTTC AGCAGTCCGG
  401  CGACGTGGTT TCGGGGCAAA GACCTGTAA ATCAGGCGGT GTTGGCGCTG
  451  TATGCGGACG AGTGGCGGCA ACCTTCGGTA CGTTGAAAA TAGGCGCAAC
  501  GTCGCACAGC CTGTGGCTCT GCACGTGCT CGGAATGCTG GTGTGCGTAT
  551  GTTGCTGCTT TTTGGTGGG CAATATACGT TCAACTGGGA AAGCACCTG
  601  TTGAGCAATG CCGCTTCGGT ACGCGCGGTG GAAATGTTGG CATGGCTGCC
  651  GTCGAAATC GGTTCCTCTG TCCCCGATGC GCGGGCGGTC ATCGAAGGCC
  701  GTCTGAACGG CAATATTGCC GATGCGCGGG CTTGGTGGG GCTGCTGGTC
  751  GGTGCTGCTG CCTGCTACGG CATCCTGCCG CGCCTGCTGG CTTGGGTAGT
  801  GTGTAAAATC CTTTGTAAAA CAAGCGAAAA CGGATTGGAT TTGGAAGAGC
  851  CCTATTATCA GGCGGTCATC CGCCGCTGGC AGAACAAAAT CACCGATGCG
  901  GATACGCGTC GGGAAACCGT GTCCGCGGTT TCACCGAAAA TCATCTTGAA
  951  CGATGCGCCG AAATGGGCGG TCATGCTGGA GACCGAGTGG CAGGACGCGC
 1001  AATGGTTCGA GGGCAGGCTG GCGCAGGAAT GGCTGGATAA GGGCGTTGCC
 1051  ACCAATCGGG AACAGGTTGC CGCGCTGGAG ACAGAGCTGA AGCAGAAACC
 1101  GGCGCAACTG CTTATCGGCG TGCGCGCCCA AACTGTGCCG GACCGCGGCG
 1151  TGTTGCGGCA GATTGTCCGA CTCTCGGAAG CGGCGCAGGG CGGCGCGGTG
 1201  GTGCAGCTTT TGGCGGAACA GGGGCTTTCA GACGACCTTT CGGAAAAGCT
 1251  GGAACATTGG CGTAACGCGC TGGCCGAATG CGGCGCGGCG TGGCTTGAGC
 1301  CTGACAGGGC GCGCAGGAA GGGCGTTTGA AAGACCAATA A

```

This corresponds to the amino acid sequence <SEQ ID 200; ORF33-1>:

```

  1  MLNPSRKLVE LVRILDEGGF IFSGDPVQAT EALRRVDGST EEKIIRRAEM
  51  IDNRNRLRET LERVAGSEFW LWVVAATFAF FTGFSVTYLL MDNQGLNFFL
 101  VLAGVLGMNT LMLAVWLAML FLRVKVGRFF SSPATWFRGK DPNVQAVLRL
 151  YADEWRQPSV RWKIGATSHS LWLCTLLGML VSVLLLLLVR QYTFNWESTL
 201  LSNAASVRV EMLAWLPSKL GFPVPDARAV IEGRLNGNIA DARAWSGLLV
 251  GSIACYGILP RLLAWVVCKI LLKTSENGLD LEKPYQAVI RRWQNKITDA
 301  DTRRETVS AV SPKII LNDAP KWA VMLETEW QDGEWFEGRL AQEWLDKGVA
 351  TNREQVAAL ETLKQKPAQL LIGVRAQTVP DRGVLRQIVR LSEAAQGGAV
 401  VQLLAEQGLS DDLSEKLEHW RNALAECEGAA WLEPDRAAQE GRLKDQ*

```

Computer analysis of this amino acid sequence gave the following results:

40 Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF33 shows 90.9% identity over a 143aa overlap with an ORF (ORF33a) from strain A of *N. meningitidis*:

```

                                     10      20      30
 45  orf33.pep                      LFLRVKVGRFFSSPATWFRXKDPVNQAVLR
                                     |||||
  orf33a      LMDNQGLNFFLVLAGVXGMNTLMLAVWLAMLFLRVKVGRFFSSPATWFRGKDPVNQAVLR
               90      100      110      120      130      140

               40      50      60      70      80      90
 50  orf33.pep      LYXDEWRXTSVRWKIXATSHSLWLCTLLGMLVSVLLLLLVRQYTFNWESTLLSNAASVRA
               || ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
  orf33a      LYADEWRXPSVRWKIGATSHSLWLCTLLGMLVSVLLLLLVRQYTFNWESTLLGDSSSVRL
               150     160     170     180     190     200

               100     110     120     130     140
 55  orf33.pep      VEMLAWLPSKLGFPVPDARSVIEGRINGNIADARAWSGLLVXSIACXGILPRL
               ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
  orf33a      VEMLAWLPAKLGFPVPDARAVIEGRINGNIADARAWSGLLVGSIACYGILPRLILAWAVCK
               210     220     230     240     250     260

 60  orf33a      ILXXTSENGLDLEKXXXXXXIRRWQNKITDADTRRETVS AVSPKIVLNDAPKWA VMLETE
               270     280     290     300     310     320

```

The complete length ORF33a nucleotide sequence <SEQ ID 201> is:

```

1   ATGTTGAATC CATCCCGAAA ACTGGTTGAG CTGGTCCGTA TTTTGAAGA
51  AGGCGGCTTT ATTTTCAGCG GCGATCCCGT GCAGGCGACG GAGGCTTTGC
101 GCGCGGTGGA CGGCAGTACG GAGGAAAAAA TCATCCGTCG GGCGAAGATG
5   151 ATCGACAGGA ACCGTATGCT GCGGGAGACG TTGGAACGTG TCGGTGCGGG
201 GTCGTTCTGG TTGTGGGTGG CGGCGGCGAC GTTTGCGTTT NTTACCGNTT
251 TTTCACTTAC TTATCTTCTA ATGGACAATC AGGGTCTGAA TTTCTTTTTG
301 GTTTTGGCGG GCGTGNTGGG CATGAATACG CTGATGCTGG CAGTATGGTT
351 GGCAATGTTG TTCCTGCGCG TGAAAGTGGG GCGTTTTTTC AGCAGTCCGG
10  401 CGACGTGGTT TCGGGGCAAA GACCTGTCA ATCAGGCGGT GTTGGCGCTG
451 TATGCGGACG AGTGGCGGCN ACCTTCGGTA CGTTGAAAA TAGGCGCAAC
501 GTCGACACAG CTGTGGCTCT GCACGCTGCT CGGAATGCTG GTGTGCGGTAT
551 TGTGCTGCTT TTTGGTGGG CAATATACGT TCAACTGGGA AAGCAGCTG
601 TTGGGCGATT CGTCTTCGGT ACGGCTGGTG GAAATGTTGG CATGGCTGCC
15  651 TGCGAAACTG GGTTTTCCCG TGCTGATGC GCGGGCGGTC ATCGAAGGTC
701 GTCTGAACGG CAATATTGCC GATGCGCGGG CTTGGTGGG GCTGCTGGTC
751 GGCAGTATCG CCTGTACGG CATCTGCCG CGCCTCTTGG CTTGGGCGGT
801 ATGCAAAATC CTTNTGNAAC CAAGCGAAAA CGGCTTGGAT TTGGAAGAGC
851 NCNNNNNTCN NNCNTCATC CGCCGCTGGC AGAACAAAAT CACCGATGCG
20  901 GATACGCGTC GGGAAACCGT GTCCGCCGTT TCGCCGAAAA TCGTCTTGAA
951 CGATGCGCGG AATGGGCGG TCATGCTGGA GACCGAATGG CAGGACGGCG
1001 AATGGTTCGA GGGCAGGCTG GCGCAGGAAT GGCTGGATAA GGGCGTTGCC
1051 GCCAATCGGG AACAGGTTGC CGCGCTGGAG ACAGAGCTGA AGCAGAAACC
1101 GGCAGCACTG CTTATCGGCG TGCGCGCCCA AACTGTGCCC GACCGCGGCG
25  1151 TGTTCGGGCA GATCGTCCGA CTTTCGGAAG CGGCGCAGGG CGGCGCGGTG
1201 GTGCANCTTT TGGCGGAACA GGGGCTTTCA GACGACCTTT CGGAAAAGCT
1251 GGAACATTGG CGTAACGCGC TGACCGAATG CGGCGCGGCG TGGCTGGAAC
1301 CCGACAGAGC GGCAGAGGAA GGCCGTCTGA AAACCAACGA CCGCACTTGA

```

This encodes a protein having amino acid sequence <SEQ ID 202>:

```

30  1   MLNPSRKLVE LVRILEEGGF IFSGDPVQAT EALRRVDGST EEKIIRRAKM
51  IDRNRMLRET LERVAGSEFW LWVAAATEAF XTXFSVTYLL MDNQGLNFFL
101 VLAGVXGMNT LMLAVWLAML FLRVKVGRRF SSPATWFRGK DPNVQAVLRL
151 YADEWRXPSV RWKIGATSHS LWLCTLLGML VSVLLLLLVLR QYTFNWESTL
201 LGDSSSVRLV EMLAWLPAKL GFPVPDARAV IEGRNLGNIA DARAWSGLLV
35  251 GSIACYGILP RLLAWAVCKI LXXTSENGLD LEKXXXXXXI RRWQNKITDA
301 DTRRETVSAP SPKIVLNDAP KWAVMLETEW QDGEWFEGRL AQEWLDKGVA
351 ANREQVAAL TELKQKPAQL LIGVRAQTVP DRGVLRQIVR LSEAAQGGAV
401 VXLLAEQGLS DDLSEKLEHW RNALTECGAA WLEPDRAAQE GRLLKNDRT*

```

ORF33a and ORF33-1 show 94.1% identity in 444 aa overlap:

```

40  10      20      30      40      50      60
    orf33a.pep  MLNPSRKLVELVRILEEGGFIFSGDPVQATEALRRVDGSTEEKIIRRAKMIDRNRMLRET
    orf33-1     MLNPSRKLVELVRILDEGGFIFSGDPVQATEALRRVDGSTEEKIIRRAEMIDRNRMLRET
100  10      20      30      40      50      60
    orf33a.pep  LERVAGSEFWLWVAAATEAFXTXFSVTYLLMDNQGLNFFLVLAGVXGMNTLMLAVWLAML
    orf33-1     LERVAGSEFWLVVAAATEAFFTGFSVTYLLMDNQGLNFFLVLAGVXGMNTLMLAVWLAML
150  70      80      90      100     110     120
    orf33a.pep  FLRVKVGRRFSSPATWFRGKDPVNQAVLRLYADEWRXPSVRWKIGATSHSLWLCTLLGML
    orf33-1     FLRVKVGRRFSSPATWFRGKDPVNQAVLRLYADEWRQPSVRWKIGATSHSLWLCTLLGML
200  130     140     150     160     170     180
    orf33a.pep  VSVLLLLLVLRQYTFNWESTLLGDSSSVRLVEMLAWLPAKLGFPVPDARAVIEGRNLGNIA
    orf33-1     VSVLLLLLVLRQYTFNWESTLLSNAASVRAVEMLAWLPKLGFPVPDARAVIEGRNLGNIA
250  190     200     210     220     230     240
    orf33a.pep  DARAWSGLLVGSIACYGILPRLLAWAVCKILXXTSENGLDLEKXXXXXXIIRWQNKITDA
    orf33-1     DARAWSGLLVGSIACYGILPRLLAWAVCKILXXTSENGLDLEKXXXXXXIIRWQNKITDA

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-158-

	orf33-1	DARAWSGLLVGSIACYGILPRLLAWVVCKILLKTSENGLDLEKPYQAVIRRWNQKITDA	250	260	270	280	290	300
5	orf33a.pep	DTRRETVSASVSPKIVLNDAPKWAVMLETEWQDGEWFEGRLAQEWLDKGVAANREQVAALF	310	320	330	340	350	360
	orf33-1	DTRRETVSASVSPKIIILNDAPKWAVMLETEWQDGEWFEGRLAQEWLDKGVAATNREQVAALF	310	320	330	340	350	360
10	orf33a.pep	TELKQKPAQLLIGVRAQTVPDGRVLRQIVRLSEAAQGGAVVXLLAEQGLSDDLSEKLEHW	370	380	390	400	410	420
	orf33-1	TELKQKPAQLLIGVRAQTVPDGRVLRQIVRLSEAAQGGAVVQLLAEQGLSDDLSEKLEHW	370	380	390	400	410	420
15	orf33a.pep	RNALTECGAAWLEPDRAAQEGRLKTNDRTX	430	440	450			
	orf33-1	RNALAECGAAWLEPDRAAQEGRLKDQX	430	440				

Homology with a predicted ORF from *N.gonorrhoeae*

ORF33 shows 91.6% identity over a 143aa overlap with a predicted ORF (ORF33.ng) from *N.gonorrhoeae*:

25	orf33.pep	LFLRVKVGRRFFSSPATWFRXKDPVNQAVLR	30
	orf33ng	LMDNQGLNFFLVLAVGLGMNTLMLAVWLATLFLRVKVGRRFFSSPATWFRGKGPVNQAVLR	100
30	orf33.pep	LYXDEWRXTSVRWKIXATSHSLWLCTLLGMLVSVLLLLLVQRQYTFNWESTLLSNAASVRA	90
	orf33ng	LYADQWRQPSVRWKIGATAHSLWLCTLLGMLVSVLLLLLVQRQYTFNWESTLLSNAASVRA	160
35	orf33.pep	VEMLAWLPSKLGFPVPDARSVIEGRNLGNIDARAWSGLLVXSIACXGILPRL	143
	orf33ng	VEMLAWLPSKLGFPVPDARAVIEGRNLGNIDARAWSGLLVGSIVCYGILPRLLAWVVCK	220

An ORF33ng nucleotide sequence <SEQ ID 203> was predicted to encode a protein having amino acid sequence <SEQ ID 204>:

40	1	MIDRDRMLRD	TLERVAGSE	WLWVVASMM	FTAGFSGTYL	LMDNQGLNFF
	51	LVLAVGLGMN	TLMLAVWLAT	LFLRVKVGRR	FSSPATWFRG	KGPVNQAVLR
	101	LYADQWRQPS	VRWKIGATAH	SLWLCTLLGM	LVSLLLLLV	ROYTFNWEST
	151	LLSNAASVRA	VEMLAWLPSK	LGFPVPDARA	VIEGRNLGNI	ADARAWSGLL
	201	VGSIVCYGIL	PRLLAWVVCK	ILLKTSENGL	DLEKTYQAV	IRRWNQKITD
	251	ADTRRETVSA	VSPKIVLND	PKWALMLETE	WQDGQWFECR	LAQEWLDKGV
45	301	AANREQVAAL	ETELKQKPAQ	LLIGVRAQTV	PDRGVLRQIV	RLSEAAQGGA
	351	VVQLLAQGL	SDDLSEKLEH	WRNALTECGA	AWLEPDRVAQ	EGRLKDQ*

Further sequence analysis revealed the following DNA sequence <SEQ ID 205>:

50	1	ATGTTGaatC	CATCCCgaAA	ACTGgttgag	ctGgTCCgtA	Ttttgaataa
	51	aggggggtTTT	attttcagcg	gcgatcctgt	gcaggcgacg	gaggctttgc
	101	gccgcgtgga	cggcAGTACG	GAggAaaaaa	tcttcctcg	GGCGGAGATg
	151	atcgACAGGg	accgtatggt	gcgggACaCg	TtggaacGTG	TGCGTGCGgg
	201	gtcgtTctgG	TTATGGGTGG	TggtggCatC	gATGATGTtt	aCCGCCGGAT
	251	TTCAGGcac	ttatCttCTG	ATGGACaatC	AGGGGCTGAA	TtTCTTTTA
	301	GTTTTggcGg	GAGTGTtggG	CATGaatacG	ctgATGCTGG	CAGTATGGtt
	351	gGCAACGTTG	TCCTGCGCG	TGAAAGTGGG	ACGGTTTTTC	AGCAGTCCGG
55	401	CGACGTGGT	TCGGGGCAAA	GGCCCTGTAA	ATCAGGCGGT	GTTGCGGCTG
	451	TATGCGGACC	AGTGGCGGCA	ACCTTCGGTA	CGATGGAAAA	TAGGCGCAAC
	501	GGCGCACAGC	TTGTGGCTCT	GCACGCTGCT	CGGAATGCTG	GTGTCGGTAT
	551	TGCTGCTGCT	TTTGGTGGCG	CAATATACGT	TCAACTGGGA	AAGCACGCTG
60	601	TTGAGCAATG	CCGCTTCGGT	ACGCGCGGTG	GAAATGTTGG	CATGGCTGCC
	651	TCGAAAAC	GTTTCCCTG	TCCCCGATGC	GCGGGCGGTC	ATCGAAGGTC
	701	GTCTGAACG	CAATATTGCC	GATGCGCGGG	CTTGCTCGGG	GCTGCTGGTC
	751	GGCAGTATCG	TCTGCTACGG	CATCCTGCCG	CGCCTCTTGG	CTTGGGTAGT

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801 GTGTAAAAATC CTTTGTGAAA CAAGCGAAAA CGGattgGAT TTGGAAAAAA
851 CCTATTATCA GGCAGTCATC CGCCGCTGGC AGAACAAAAT CACCGATGCG
901 GATACGCGTC GGGAAACCGT GTCCGCGGTT TCGCcgAAAA TCGTCTTGAA
951 CGATGCGCCG AAATGGGCGC TCATGCTGGA GACCGAGTGG CAGGACGGCC
1001 AATGGTTCGA GGGCAGGCTG GCGCAGGAAT GGCTGGATAA GGGCGTTGCC
1051 GCCAATCGGG AACAGGTTGC CGCGCTGGAG ACAGAGCTGA AGCAGAAACC
1101 GGCACCACTG CTTATCGGCG TACGCGCCCA AACTGTGCCG GACCGGGGCG
1151 TGCTGCGGCA GATTGTGCGG CTTTCGGAAG CGGCGCAGGG CGGCGCGGTG
1201 GTGCAGCTTT TGGCGGAACA GGGGCTTTCA GACGACCTTT CGGAAAAAGCT
1251 GGAACATTGG CGTAACGCGC TGACCGAATG CGGCGCGGCG TGGCTTGAGC
1301 CTGACAGGGT GCGCGAGGAA GGCCGTTTGA AAGACCAATA A

```

This encodes a protein having amino acid sequence <SEQ ID 206; ORF33ng-1>:

15
20
25
30
35
40
45
50
55
60
65

```

1 MLNPSRKLVE LVRILNKGFF IFSGDPVQAT EALRRVDGST EEKIFRRAEM
51 IDRDRMLRDT LERVAGSFW LWVVVASMFF TAGFSGTYLL MDNQGLNFFL
101 VLAGVLGMNT LMLAVWLATL FLRVKVGRRF SSPATWFRGK GPVNQAVLRL
151 YADQWRQPSV RWKIGATAHS LWLCTLLGML VSVLLLLLVLR QYTFNWESTL
201 LSNAASVRV EMLAWLPSKL GFPVPDARAV IEGRNLGNIA DARAWSGLLV
251 GSIVCYGILP RLLAWVVKI LLKTSENGLD LEKTYQYQAVI RRWQNKITDA
301 DTRRETSAV SPKIVLNDAP KWALMLETW QDQWFEGR LAQEWLDKGVA
351 ANREQVAAL TELKQKPAQL LIGVRAQTVP DRGVLRQIVR LSEAAQGGAV
401 VQLLAEQGLS DDLSEKLEHW RNALTECGAA WLEPDRVAQE GRLKDQ*

```

ORF33ng-1 and ORF33-1 show 94.6% identity in 446 aa overlap:

25
30
35
40
45
50
55
60
65

```

      10      20      30      40      50      60
orf33-1.pep MLNPSRKLVELVRILDEGGFIFSGDPVQATEALRRVDGSTEEKIIRRAEMIDNRMLRET
      10      20      30      40      50      60
orf33ng-1   MLNPSRKLVELVRILNKGFFIFSGDPVQATEALRRVDGSTEEKIFRRAEMIDRDRMLRDT

      70      80      90     100     110     120
orf33-1.pep LERVAGSFWLWVVAATFAFFTGFSVTYLLMDNQGLNFFLVLAGVLGMNTLMLAVWLAML
      70      80      90     100     110     120
orf33ng-1   LERVAGSFWLWVVVASMFFTAGFSGTYLLMDNQGLNFFLVLAGVLGMNTLMLAVWLATL

      130     140     150     160     170     180
orf33-1.pep FLRVKVGRRFSSPATWFRGKDPVNQAVLRRLYADEWRQPSVRWKIGATSHSLWLCTLLGML
      130     140     150     160     170     180
orf33ng-1   FLRVKVGRRFSSPATWFRGKGPVNQAVLRRLYADQWRQPSVRWKIGATAHSLWLCTLLGML

      190     200     210     220     230     240
orf33-1.pep VSVLLLLLVQRQYTFNWESTLLSNAASVRVEMLAWLPSKLGFPVPDARAVIEGRNLGNIA
      190     200     210     220     230     240
orf33ng-1   VSVLLLLLVQRQYTFNWESTLLSNAASVRVEMLAWLPSKLGFPVPDARAVIEGRNLGNIA

      250     260     270     280     290     300
orf33-1.pep DARAWSGLLVGSIACYGILPRLAWVVKILLKTSENGLDLEKPYQYQAVIRRWQNKITDA
      250     260     270     280     290     300
orf33ng-1   DARAWSGLLVGSIACYGILPRLAWVVKILLKTSENGLDLEKTYQYQAVIRRWQNKITDA

      310     320     330     340     350     360
orf33-1.pep DTRRETSAVSPKIIINDAPKWAVMLETWQDGEWFEGR LAQEWLDKGVA TNREQVAAL
      310     320     330     340     350     360
orf33ng-1   DTRRETSAVSPKIVLNDAPKWALMLETWQDGEWFEGR LAQEWLDKGVA ANREQVAAL

      370     380     390     400     410     420
orf33-1.pep TELKQKPAQLLIGVRAQTVPDRGVLRQIVRLSEAAQGGAVVQLLAEQGLSDDLSEKLEHW
      370     380     390     400     410     420
orf33ng-1   TELKQKPAQLLIGVRAQTVPDRGVLRQIVRLSEAAQGGAVVQLLAEQGLSDDLSEKLEHW

      430     440
orf33-1.pep RNALAECCGAAWLEPDRAAQEGRLKDQX
      430     440
orf33ng-1   RNALAECCGAAWLEPDRAAQEGRLKDQX

```

Example 25

10	1	..CAGAAGAGTT	TGTCGAGAAT	TTCTTTATGG	GGTTTGGGCG	GCGTGTTTTT
	51	CGGGGTGTCC	GGTCTGGTAT	GGTTTTCTTT	GGGCGTTTCT	TT.GAGTGCG
	101	CCTGTTTTTC	GGGTGTTTCT	TTTCGGGGTT	CGGGACGGGG	GACGTTTGTG
	151	GGCAGTACGG	GGGTTTCTTT	GAGTGTGTTT	TCAGCTTGTG	TTCC.GGCGT
	201	CGTCCGGCTG	CCTGTCGGTT	TGAGCTGTGT	CGCGAGGTTG	G..GTTTGA
	251	CCCGGTTTTT	CTTGGGTGCG	GCAGGGGACG	TCATTCTCCT	CCCCTTTTCG
15	301	TCTGTGCCGT	CCGGCTGTGC	GGGTTCGGAT	GAGGCGGCGT	GGTGGTGTTC
	351	GGGTTGGGCG	GCATCTTGTT	CCGACTACGC	CGTTTGGCAG	CCAGAATTTCG
	401	GTTCGCGGG	GGCTGTCGGT	GTGTTGCGGT	TCGGCTTGAA	GGGTTTTTGT
	451	GTCC..				

20 1 ..QKSLSRISLW GLGGVFFGVG GLVWFSLGVS XECACFSGVS FRGSGRGTFV
51 GSTGVSLSVF SACVXGVVRL PVGLSCVGRL XXLTRFLGA AGDVILLPLS
101 SVPGCGAGSD EAAWWCSGWA ASCPTTPFGS QNSVSRGLSV CCGSA*RVLS
151 S...

25	1	ATGATGATGC	CGTTCATAAT	GCTTCCTTGG	ATTGCKGGTG	TGCCTGCCGT
	51	GCCGGGTCAG	AATAGGTTGT	CCAGAATTTC	TTTATGGGGT	TTGGGCGGCG
	101	TGTTTTTCGG	GGTGTCCGGT	TTGGTATGGT	TTTCTTTGGG	CGTTTTCTTTG
	151	GGCTGCGCCT	GTTTTTCGGG	TGTTTTCTTT	CGGGGTTFCG	GACGGGGGAC
30	201	GTTTGTGGCG	AGTACGGGGG	TTTCTTTGAG	TGTTCTTTTCA	GCTTGTGTTTC
	251	CGGCGTCGTC	CGGCTGCCTG	TCGGTTTGAG	CTGTGTCCGC	AGGTTGCGGT
	301	TTGACCCGGT	TTTTCTTTGG	TGCGGCAGGG	GACGGCAGTC	CGCTGCCGCT
	351	TTGCTCTGTG	CCGTCGGGCT	GTGCGGGTTC	GGATGAGGCG	CGCTGGTGGT
35	401	GTTCCGGTTG	GCGCGCATCT	TGTCCGACTA	CGCGGTTTGG	CAGCCAGAAT
	451	TCGGTTTCGC	GGGGGCTGTC	GGTGTGTTGC	GGTTCGGCTT	GAAGGGTTTT
	501	GTCGCGGTTT	GGGTTGAATG	TGCTGACGAT	GCCTATTGCC	AATGCGCCGA
	551	TGGCGGCGAT	ACAGATGAGC	AATACGGCGC	GTTATCAGGAG	TTTTGGGGTC
40	601	AGCCTGAAGG	GTTTGTTCGG	TTTTTTTGCC	ATTTTGTATTG	TGTTTTTTGGG
	651	GTGTCGGGCA	ATGCCGTCTG	AAGGCGGTTT	AGACGGCATT	GCCGAGTCAG
	701	CGTTGGACGT	AGTTTTTGTA	GAGGGTGATG	ACTTTTTGTA	CGCCGACGGT
	751	GGTGCTGACT	TTTTGGGTAA	TCTGCGCTTG	TCTCTCGGGG	GTCAGGATGC
45	801	CCATAACGTA	GGTTACGTTG	CCGTAGGTAA	CGATTTTGAC	GCGCGCCTGT
	851	GTGGCGGGGC	TGATGCCCAA	CAGCGTGGCG	CGGACTTTGG	ATGTGTTCCA
	901	AGTGTCGCCG	GCGATGTCGC	CGGCAGTGCG	CGGCAGGGAG	GCGACGGTAA
	951	TATAGTTGTA	CACGCCTTCG	GCGGCCTGTT	CGGAACGTGC	AATCTGACCG
50	1001	ACGAACGTGT	TTTCGCCTTC	GGTGGCGACT	TGTCCGAGCA	GCAGCAGGTG
	1051	GCGGTTGTAG	CCGACGACGG	AGATTTGGGG	CGTGTAGCCT	TTGGTTTGGT
	1101	TGTTTTGGCG	CAGATAGGAA	CGGGCGGTGT	TTTCGATACG	CAACGCCATA
	1151	ACGTTGTGCT	GCGTTTGC GC	GCCGTTGGTT	GCGCGGTGCA	CGGCGGATTT
55	1201	GCGGCCGACG	GCGGCGCTTC	CGATTACTGC	GCTGACGCAG	CCGCTAAGGG
	1251	CAAGGCTGAA	AATGGCGGCA	ATCAGGCTGC	GGACGGTGTG	CGGTTTGGGT
	1301	TTCATCCGGT	GCTTCCTTTC	TTGGGCGTTT	CAGACGGCAT	TGCTTTGCGC
	1351	CATGCCGTCT	GA			

55

1	<u>MMMPFIMLPW</u>	<u>IAGVPAVFGQ</u>	<u>NRLSRISLWG</u>	<u>LGGVFFGVSG</u>	<u>LWVFSLGVSL</u>
51	<u>GCACFSGVSF</u>	<u>RGSGRGTFVG</u>	<u>STGVSLSVFS</u>	<u>ACVPASSGCL</u>	<u>SV*AVSAGCG</u>
101	<u>LTRFFLGAAG</u>	<u>DGSPPLSSV</u>	<u>PSGCAGSDEA</u>	<u>AWWCSGWAAS</u>	<u>CPTTFPGSQN</u>
151	<u>SVSRGLSVCC</u>	<u>GSA*RVLSPF</u>	<u>GLNVLTMPIA</u>	<u>NAPMAAIQMS</u>	<u>NTARIRSLGV</u>

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201 SLKGLFGFFA ILIVLLGCRA MPSEGGSDGI AESALDVVLV EGDDFLYADG
 251 GADFLGNLRL FFGGEDAHNV GYVAVGNDFD ARLCGGADAQ QRGADFGCVP
 301 SVAGDVAGSA RQGGDGNIVV HAFGGLFGTC NLTDELFFAF GDDLSEQQQV
 351 AVVADDGDLG RVAFGILVVLA QIGTGGGFDT QRHNVVVGLR AGGSAVDGGF
 5 RADGGASDYC ADAAAKGKAE NGGNQGADGV RFGFHRVLPF LGVSDGIALR
 451 HAV*

Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF34 shows 73.3% identity over a 161aa overlap with an ORF (ORF34a) from strain A of *N.*

10 *meningitidis*:

```

                                10      20      30
orf34.pep      QKSLSRISLWGLGGVFFGVSGLVWFSGLVGSXE-----CAC
                || ||| ||||| ||||| ||||| ||||| ||||| |||||
orf34a      MMXFXIMLPWIAGVPAVPGQKRLSRXSLWGLGGXFFGVSGLVWFSGLVGSXSXSLGVSXGCAC
                10      20      30      40      50      60

                                40      50      60      70      80      90
orf34.pep      FSGVSRGSGRGTFFVGSTGVSLSVFSACVXGVVRLPVGLSCVGRLLX-----LTRFFLGA
                ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
orf34a      FSGVSRGSGRGTFFVGSTGVSLSVFSACA-----PASSGCLSVXAVSAGCGLTRFXFGA
                70      80      90      100      110

                                100      110      120      130      140      150
orf34.pep      AGDVILLPLSSVPSGCAGSDEAAWNCSGWAASCPTTFPGSQNSVSRGLSVCCGSAXRVLS
                ||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
orf34a      AGDGSPLPLSSVPSGCAGADEEAXXCSGWAASCPTTFPGSQNSVSRGLSVCCGSVWRVLS
                120      130      140      150      160      170

                                180      190      200      210      220      230
orf34.pep      S
orf34a      PFGXNVLTMPIANAPMAVIQMSNTARIRSLGVSLKGLFXFFAILIVLLGCRAMPSEGGSD

```

The complete length ORF34a nucleotide sequence <SEQ ID 211> is:

```

35      1  ATGATGATNC  CGTTNATAAT  GCTTCCTTGG  ATTGCGGGTG  TGCCTGCCGT
      51  GCCGGGTCAG  AAGAGGTTGT  CGAGAANTTC  TTTATGGGGT  TTAGGCGGCN
     101  TGTTTTTCGG  GGTGTCCGGT  TTGGTATGGT  TTTCTTTGGG  CGTTTCTNTT
     151  TCTTTGGGTG  TTTCTNTGGG  CTGTGCCTGT  TTTTCGGGTG  TTTCTTTTCG
     201  GGGTTCGGGA  CGGGGGACGT  TTGTGGGCAG  TACNCGGGTT  TCTTTGAGTG
     40  251  TGTTTTCAGC  TTGTGCTCCG  GCGTCGTCCG  GCTGCCTGTC  GGTTTNAGCT
     301  GTGTCCGCAG  GTTGCGGTTT  GACCCGNTT  TTCTTNGGTG  CGGCAGGGGA
     351  CGGCAGTCCG  CTGCCGCTTT  CGTCTGTGCC  GTCCGGCTGT  GCGGGTGCGG
     401  ATGAGGAGGC  GTNGTNGTGT  TCGGGTTGGG  CGGCATCTTG  TCCGACTACG
     45  451  CCGTTTGGCA  GCCAGAATTC  GGTTTCGCGG  GGGCTGTCCG  TGTGTTGCGG
     501  TTCGGTNTGG  AGGGTTTGT  CNCCGTTCGG  GTNGAATGTG  CTGACGATGC
     551  CTATTGCCAA  TGCGCCGATG  GCGGTGATAC  AGATAGACAA  TACGGCGCGT
     601  ATCAGGAGTT  TGGGGGTCAG  CCTGAAGGGT  TTGTTCTNGT  TTTTGGCCAT
     651  TTTGATTGTG  CTTTGGGGT  GTCGGGCAAT  GCCGTCTGAA  GCGGGTTCAG
     701  ACGGCATTGC  CGAGTCAGCG  TTGGACGTAG  TTTNGGTAGA  GGGTGATGAC
     50  751  TTTTGTACG  CCGACGGTGG  TGCTGACTTT  TTGGGTAATC  TGCGCCTGTT
     801  CTTCCGGGGT  GAGGATGCC  ATAACGTAGG  TTACGTTGCC  GTAGGTAACG
     851  ATTTTGACGC  GCGCCTGTGT  GCGGGGCTG  ATGCCCAACA  GCGTGGCGCG
     901  GACTTTGGAT  GTGTTCCAAG  TGTGCGCGGC  GATGTCGCCG  GCAGTGCGCG
     951  GCAGGGAGGC  GACGGTAATG  TANTTGTACA  CGCCTTCGGC  GGCCTGTTCG
     55  1001  GAACGTGCAA  TCTGACCAGC  GAACTGTTTC  TCGCCTTCGG  TGGCGACTTG
     1051  TCCGAGCAGC  AGCAGGTGGC  GGTGTAGGCC  GACAACGGAG  ATTTGGGCGC
     1101  TGTANCTTT  GGTGTTGGTG  TTTTGGCGCA  GATAGGAGCG  GCGGTGGTGT
     1151  TCGATACGCA  GCGCCATTAC  GTGTGCTGTC  GTNCGCGCG  CGGTGGTTCG
     1201  GCGGTCGACG  GCGGATTTCG  CGCCGACCGC  CGCGCCGCCG  ACGACTGCGC
     60  1251  TGACGCAGCC  GCGAGGGGCA  AGGCTGAGGA  CGGCGGCAGT  CAGGGTGCGG
     1301  ACGGTGTGCG  GTTTGGGTTT  CATCGGGTGC  TTCCTTTCTT  GGGCGTTTCA
     1351  GACGGCATTG  CTTTGCGCCA  TGCCGTCTGA

```

This encodes a protein having amino acid sequence <SEQ ID 212>:

```

1  MMXPXIMLPW IAGVPAVPGQ KRLSRXSLWG LGGXFFGVSG LVWFSLVGSX
51  SLGVSXGCAC FSGVSFRGSG RGTfVGSTGV SLSVFSACAP ASSGCLSVXA
101 VSAGCGLTRX FXGAAGDGSP LPLSSVPSGC AGADEEAXXC SGWAASCPTT
151 PFQSQNSVSR GLSVCCGSVW RVLSPFGXNV LTMPIANAPM AVIQMSNTAR
201 IRSLGVSLKG LEXFFAILIV LLGCRAMPSE GGSDDGIAESA LDVVXVEGDD
251 FLYADGGADF LGNLRLEFFGG EDAHNVGYYA VGNDFDARLC GGADAQQRGA
301 DFGCVPSVAG DVAGSARQGG DGNVXVHAFG GLFGTCNLTD ELFLAFGGDL
351 SEQQQVAVVA DNGDLGRVXF GLVVLAQIGA GGGFDTQRHY VVVGXRAGGS
401 AVDGGFRADR RAADDCADAA AEGKAEDGGS QGADGVREGF HRVLPFLGV
451 DGIALRHAV*

```

ORF34a and ORF34-1 show 91.3% identity in 459 aa overlap:

```

15  orf34a.pep      10      20      30      40      50      60
      MMXPXIMLPW IAGVPAVPGQ KRLSRXSLWG LGGXFFGVSG LVWFSLVGSX SLSGV
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
orf34-1  MMMPFIMLPW IAGVPAVPGQ NRLSRISLWGLGGVFFGVSG LVWFSLVGSX -----GCAC
      10      20      30      40      50

20  orf34a.pep      70      80      90      100     110     120
      FSGVSFRGSG RGTfVGSTGV SLSVFSACAP ASSGCLSVXAVSAGCGLTRXFXGAAGDGSP
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
orf34-1  FSGVSFRGSG RGTfVGSTGV SLSVFSACV PASSGCLSVXAVSAGCGLTRFFLGAAGDGSP
      60      70      80      90      100     110

25  orf34a.pep      130     140     150     160     170     180
      LPLSSVPSGCAGADEEAXXC SGWAASCPTTPFGSQNSVSRGLSVCCGSVWRVLSPFGXNV
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
orf34-1  LPLSSVPSGCAGSDEAAW CSGWAASCPTTPFGSQNSVSRGLSVCCGSAXRVLSPFGLNV
      120     130     140     150     160     170

30  orf34a.pep      190     200     210     220     230     240
      LTMPIANAPMAVIQMSNTARIRSLGVSLKGLFXFFAILIVLLGCRAMPSEGGSDGIAESA
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
orf34-1  LTMPIANAPMAAIQMSNTARIRSLGVSLKGLFGFFAILIVLLGCRAMPSEGGSDGIAESA
      180     190     200     210     220     230

35  orf34a.pep      250     260     270     280     290     300
      LDVVXVEGDDFLYADGGADFLGNLRLEFFGGEDAHNVGYYAVGNDFDARLCGGADAQQRGA
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
orf34-1  LDVVLVEGDDFLYADGGADFLGNLRLEFFGGEDAHNVGYYAVGNDFDARLCGGADAQQRGA
      240     250     260     270     280     290

40  orf34a.pep      310     320     330     340     350     360
      DFGCVPSVAGDVAGSARQGGDGNVXVHAFGGLFGTCNLTD ELFLAFGGDLSEQQQVAVVA
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
orf34-1  DFGCVPSVAGDVAGSARQGGDGNIVVHAFGGLFGTCNLTD ELFFAFGGDLSEQQQVAVVA
      300     310     320     330     340     350

45  orf34a.pep      370     380     390     400     410     420
      DNGDLGRVXFGLVVLAQIGAGGGFDTQRHYVVVGXRAGGS AVDGGFRADRRRAADDCADAA
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
orf34-1  DDGDLGRVAFGLVVLAQIGTGGGFDTQRHNVVVGRLAGGS AVDGGFRADGGASDYCADAA
      360     370     380     390     400     410

50  orf34a.pep      430     440     450     460
      AEGKAEDGGSQGADGVRFGRVLPFLGVSDGIALRHAVX
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
orf34-1  AKGKAENGNGQADGVRFGRVLPFLGVSDGIALRHAVX
      420     430     440     450

```

Homology with a predicted ORF from *N.gonorrhoeae*

ORF34 shows 77.6% identity over a 161aa overlap with a predicted ORF (ORF34.ng) from *N. gonorrhoeae*:

```

orf34.pep      QKLSLRISLWGLGGVFFGVSG LVWFSLVGSX -----CAC      35

```


	orf34ng	MMMPFIMLPWIAGVPAVPGQKRLSRISLWGLAGVFFGVSGLVWFSLVGSFSLGVSLGCAC	60
	orf34.pep	FSGVSFRGSGRGTFVGGSTGVSLSVFSACVXGVVRLPVGLSCV-----GRLXXLTRFFLGA	90
5	orf34ng	FSGVSFRGSGWGAFAVGSTGVSLSVFSACVP----VPVNESAARAASEGR--GLTRFFLGA	114
	orf34.pep	AGDVILLPLSSVPSGCAGSDEAAWCSGWAASCPTTFPGSQNSVSRGLSVCCGSAXRVLS	150
10	orf34ng	AGDGSPLPLSSVPSGCAGSDEAAWCSGWAASCPTAPFGSQNSVSRGLSVCCGSVWRVLS	174
	orf34.pep	S	175
	orf34ng	PFGLNVLTMPTANAPMAVIQMSNTARIRSLGVSLKGLFGFFAILIVLLGCRAMPSEGGSD	234

15 The complete length ORF34ng nucleotide sequence <SEQ ID 213> is:

	1	ATGATGATGC	CGTTCATAAT	GCTTCCTTGG	ATTGCGGGTG	TGCCTGCCGT
	51	GCCGGGTCAA	AAGAGGTTGT	CGAGAATCTC	TTTATGGGGT	TTGGCCGGCG
	101	TGTTTTTCGG	GGTGTCGGGT	TTGGTATGGT	TTTCTTTGGG	CGTTTCTTTT
	151	TCTTTGGGTG	TTTCTTTGGG	CTGCCCTGT	TTTTCGGGTG	TTTCTTTTCG
20	201	GGGTTCGGGA	TGGGGGGCGT	TTGTGGGCAG	TACGGGGGTT	TCTTTGAGTG
	251	TGTTTTTCAGC	TTGTGTTCGG	GTGCCGGTTA	ACGAATCGGC	TGCCCCGGGC
	301	GCATCCGAAG	GGCGCGGTTT	gACCCGGTTT	TTCTTGGGTG	CGGCAGGGGA
	351	CGGCAGTCCG	CTGCCGCTTT	CTTCTGTGCC	GTCCGGCTGT	GCGGGTTCGG
	401	ATGAGGCGGC	GTGGTGGTGT	TCCGGTTGGG	CGGCATCTTG	TCCGACGGCG
25	451	CCGTTTGGCA	GCCAGAATTC	GGTTCGCGCG	GGGCTGTCCG	TGTGTGCGCG
	501	TTCCGTTTGG	AGGGTTTTGT	CGCCGTTCCG	GTTGAATCTG	CTGACGATGC
	551	CTACTGCCAA	TGCGCCGATG	GCGGTGATAC	AGATGAGCAA	TACGGCGCGT
	601	ATCAGGAGTT	TGGGGGTGAG	CCTGAAGGGT	TTGTTGCGTT	TTTTTGCCAT
	651	TTTGATTGTG	CTTTTGGGGT	GTCCGGCAAT	GCCGCTGTAA	GGCGGTTTCT
30	701	ACGGCATTGC	CGAGTCAGCG	TTGGACGTAG	TTTTGGTAGA	GGTAATGAC
	751	TTTTTGTAAC	CCGACggtGG	TGCTGACTTT	TTGGGTAATC	TGCGGCTGTT
	801	CTTCGGGGGT	GAGGATGCCC	ATAACGTAGG	TTACATTGCC	GATAGTAATG
	851	ATTTTGACGC	GCGCCTGTGT	AGCGGGGCTG	ATGCCCAGCA	GcgtgCGCGC
	901	GACTTTGGAC	GTGTTCCAAG	TGTCGCCGGC	GATGTCGCCC	GCAGTGCGCG
35	951	GCAGGGAGGC	GACGGTAATG	TAGTTGTATA	CGCCTTCGGC	GGCCTGTTCC
	1001	GAACGTGCAA	TCTGACCGAC	GAACGTTTTT	TGCGCTTCGG	TGGCGACTTG
	1051	TCCGAGCAGC	AGCAGGTGGC	GGTTGTAGCC	GACGACGGAG	ATTTGGGGCG
	1101	TGTAGCCTTT	GGTTTGGTTG	TTTTGGCGCA	GGTAGGAACG	GCGGTGGTT
	1151	TCGATACGCA	ACGCCATAAC	GTtgtCATCG	GTTtgtcgcg	CGGTGGTTcg
40	1201	gCGGTCGATG	ACGGATTTTG	CGCCGACGGC	GGCCCGCCCG	ACGACTGCGC
	1251	TGAAGCAGCC	GCCGAGGGCA	AGGCTGAGGA	CGGCGGCAAT	CAGGGTGCGG
	1301	ACGGTGTGTG	GTTTGGGTTT	CATCGGGGAC	TTCCTTTCTT	GGGCGTTTCA
	1351	GACGGCATTG	CTTTGCGCCA	TGCCGTCTGA		

This encodes a protein having amino acid sequence <SEQ ID 214>:

45	1	MMMPFIMLPW	IAGVPAVPGQ	KRLSRISLWG	LAGVFFGVSG	LVWFSLVGSF
	51	SLGVSLGCAC	FSGVSFRGSG	WGAFAVGSTGV	SLSVSFACVP	VPVNESAARA
	101	ASEGRGLTRF	FLGAAGDGSP	LPLSSVPSGC	AGSDEAAWVC	SGWAASCPTA
	151	PFGSQNSVSR	GLSVCCGSVW	RVLSPFGLNV	LTMPTANAPM	AVIQMSNTAR
50	201	IRSLGVSLKG	LFGFFAILIV	LLGCRAMPSE	GGSDGIAESA	LDVVLVGEND
	251	FLYADGGADF	LCNLRLFFGG	EDAHNVGYIA	VGNDFDARLC	SGADAQQRGA
	301	DFGRVPSVAG	DVARSAHQGG	DGNVVVYAFG	GLFGTCNLTD	ELFFAFGGDL
	351	SEQQQVAVVA	DDGDLGRVAF	GLVVLAQVGT	GGGFDTQRHN	VVIGLRAGGS
	401	AVDDGFCADG	GPADDCAEAA	AEGKAEDGGN	QGADGVWFGF	HRGLPFGLVS
	451	DGIALRHAV*				

55 ORF34ng and ORF34-1 show 90.0% identity in 459 aa overlap:

		10	20	30	40	4	50
	orf34-1.pep	MMMPFIMLPWIAGVPAVPGQNRSLSRISLWGLGGVFFGVSGLVWFSLVGS-----LGCAC					
	orf34ng	MMMPFIMLPWIAGVPAVPGQKRLSRISLWGLAGVFFGVSGLVWFSLVGSFSLGVSLGCAC					
60		10	20	30	40	50	60
	orf34-1.pep	FSGVSFRGSGRGTFVGGSTGVSLSVFSACVPASSGCLSVXAVSAGCGLTRFFLGAAGDGSP					
	orf34ng	FSGVSFRGSGWGAFAVGSTGVSLSVFSACVPVPVNESAARAASEGRGLTRFFLGAAGDGSP					

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		70	80	90	100	110	120
		120	130	140	150	160	170
5	orf34-1.pep	LPLSSVP	SGCAGS	DEAAW	CSGWA	ASCPTT	PFQSQNSVSRGLSVCCGSAXRVLSPFGLNV
	orf34ng	LPLSSVP	SGCAGS	DEAAW	CSGWA	ASCPTT	PFQSQNSVSRGLSVCCGSVWRVLSPFGLNV
		130	140	150	160	170	180
		180	190	200	210	220	230
10	orf34-1.pep	LTMPIAN	APMAAI	QMSNTA	RIRSLG	VSLKGL	FGFFAILIVLLGCRAMPSEGGSDGIAESA
	orf34ng	LTMPAN	APMAVI	QMSNTA	RIRSLG	VSLKGL	FGFFAILIVLLGCRAMPSEGGSDGIAESA
		190	200	210	220	230	240
		240	250	260	270	280	290
15	orf34-1.pep	LDVVLVE	GDDFLY	ADGGAD	FLGNLRL	FFGGEDA	HNVGYVAVGNDFDARLCGGADAQQRGA
	orf34ng	LDVVLVE	GDDFLY	ADGGAD	FLGNLRL	FFGGEDA	HNVGYVAVGNDFDARLCGGADAQQRGA
		250	260	270	280	290	300
		300	310	320	330	340	350
20	orf34-1.pep	DFGCVPS	VAGDVA	GSARQQ	GGDGNIV	VHAFGGL	FGTCNLTDDELFFAFGGDLSEQQQVAVVA
	orf34ng	DFGRVPS	VAGDVARS	ARQQGG	DGNVVV	YAFGGL	FGTCNLTDDELFFAFGGDLSEQQQVAVVA
25		310	320	330	340	350	360
		360	370	380	390	400	410
30	orf34-1.pep	DDGDLGR	VAFGLV	VLAQIG	TGGGFD	TQRHN	VVVGRLRAGGS
	orf34ng	DDGDLGR	VAFGLV	VLAQIG	TGGGFD	TQRHN	VVVGRLRAGGS
		370	380	390	400	410	420
		420	430	440	450		
35	orf34-1.pep	AKGKAEN	GGNQGA	DGVRFG	FHRVLP	FLGVSD	GIALRHAVX
	orf34ng	AEGKAED	GGNQGA	DGVWFG	FHRGLP	FLGVSD	GIALRHAVX
		430	440	450	460		

Based on this analysis, including the presence of a putative leader sequence (double-underlined) and several putative transmembrane domains (single-underlined) in the gonococcal protein, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 26

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 215>:

45	1	ATGAAAACCT	TCTTCAAAC	CCTTTCCGCC	GCCGCACTCG	CGCTCATCCT
	51	CGCCGCCTGC	GGATT.CAAA	AAGACAGCGC	GCCCGCCGCA	TCCGCTTCTG
	101	CCGCCGCCGA	CAACGGCGCG	GCGTAAAAAA	GAAATCGTCT	TCGGCACGAC
	151	CGTCGGCGAC	TTCGGCGATA	TGGTCAAAGA	ACAAATCCAA	GCCGAGCTGG
	201	AGAAAAAGG	CTACACCGTC	AAACTGGTCG	AGTTTACCGA	CTATGTACGC
	251	CCGAATCTGG	CATTGGCTGA	GGCGAGTTG		

50 This corresponds to the amino acid sequence <SEQ ID 216; ORF4>:

	1	MKTFFKTL	SA AALALIL	AAAC G.QKDS	APAA SASAA	ADNGA AKKEIV	FGTT
	51	VGDFGDM	VKE QIQAE	LEKKG YTVKL	VEFTD YVRPN	LALAE GEL	

Further sequence analysis revealed the complete nucleotide sequence <SEQ ID 217>:

55	1	ATGAAAACCT	TCTTCAAAC	CCTTTCCGCC	GCCGCACTCG	CGCTCATCCT
	51	CGCCGCCTGC	GGCGGTCAA	AAGACAGCGC	GCCCGCCGCA	TCCGCTTCTG
	101	CCGCCGCCGA	CAACGGCGCG	GCGAAAAAAG	AAATCGTCTT	CGGCACGACC
	151	GTCGCGGACT	TCGGCGATAT	GGTCAAAGAA	CAAATCCAAG	CCGAGCTGGA
	201	GAAAAAAGGC	TACACCGTCA	AACTGGTCGA	GTTTACCGAC	TATGTACGCC

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251 CGAATCTGGC ATTGGCTGAG GCGAGTTGG ACATCAACGT CTTCCAACAC
301 AAACCCCTATC TTGACGACTT CAAAAAAGAA CACAATCTGG ACATCACCGA
351 AGTCTTCCAA GTGCCGACCG CGCCTTTGGG ACTGTACCCG GGCAAGCTGA
401 AATCGCTGGA AGAAGTCAAA GACGGCAGCA CCGTATCCGC GCCCAACGAC
5 451 CCGTCCAACT TCGCCCGCGT CTGGTGATG CTCGACGAAC TGGGTTGGAT
501 CAAACTCAAA GACGGCATCA ATCCGTGAC CGCATCCAA GCGGACATCG
551 CCGAGAACCT GAAAAACATC AAAATCGTCG AGCTTGAAGC CGCGCAACTG
601 CCGCGTAGCC GCGCCGACGT GGATTTTGCC GTCGTCAACG GCAACTACGC
651 CATAAGCAGC GGCATGAAGC TGACCGAAGC CCTGTTCCAA GAACCGAGCT
10 701 TTGCCTATGT CAACTGGTCT GCCGTCAAAA CCGCCGACAA AGACAGCCAA
751 TGGCTTAAAG ACGTAACCGA GGCCTATAAC TCCGACGCGT TCAAAGCCTA
801 CGCGCACAAA CGCTTCGAGG GCTACAAATC CCCTGCCGCA TGAATGAAG
851 GCGCAGCCAA ATAA

```

This corresponds to the amino acid sequence <SEQ ID 218; ORF4-1>:

```

15 1 MKTFFKTLSA AALALILAAAC GGQKDSAPAA SASAAADNGA AKKEIVFGTT
51 VGDFGDMVKE QIQAELEKKG YTVKLVEFTD YVRPNLALAE GELDINVFQH
101 KEYLDDEFKE HNLDTVEVFQ VETAPLGLYP GKLSLEEVK DGSTVSAPND
151 PSNFARVLVM LDELGWIKLK DGINPLTASK ADIAENLKNI KIVELEAAQL
201 PRSRADVDFV VVNGNYAISS GMKLTEALFQ EPSFAYVNW AVKTADKDSQ
20 251 WLKDVTEAYN SDAFKAYAHK RFEGYKSPAA WNEGAAK*

```

Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF4 shows 93.5% identity over a 93aa overlap with an ORF (ORF4a) from strain A of *N.*

meningitidis:

```

25 orf4.pep      10      20      30      40      50      59
      MKTFFKTLSAALALILAAACG-QKDSAPAASASAAADNGAAKKEIVFGTTVGDFGDMVKE
      |||||
orf4a      MKTFFKTLSAALALILAAACGGQKDSAPAASASAAADNGAAKKEIVFGTTVGDFGDMVKE
      10      20      30      40      50      60

30 orf4.pep      60      70      80      90
      QIQAELEKKG YTVKLVEFTD YVRPNLALAE GEL
      || |||||
orf4a      XIQPELEKKG YTVKLVEFTD YVRPNLALAE GELDINXQHXXYLDDXKXHNLDITXVXQ
      70      80      90      100     110     120

35 orf4a      VPTAPLGLYPGKLSLXXVKXGSTVSAPNDPXXFXRVLVMLDELGXIKLKDXIXXXXXXX
      130     140     150     160     170     180

```

The complete length ORF4a nucleotide sequence <SEQ ID 219> is:

```

40 1 ATGAAACCT TCTTCAAAAC CCTTCCGCC GCCGCACTCG CGCTCATCCT
51 CGCCGCCTGC GGCGGTCAAA AAGATAGCGC GCCCGCCGCA TCCGCTPTCTG
101 CCGCCGCCGA CAACGGCGCG GCGAANAAAG AAATCGTCTT CGGCACGACC
151 GTCGGCGACT TCGGCGATAT GGTCAAAGAA CANATCCAAC CCGAGCTGGA
201 GAAAAAAGGC TACACCGTCA AACTGGTCTGA GTNTACCGAC TATGTGCGCN
45 251 CGAATCTGGC ATTGGCTGAG GCGAGTTGG ACATCAACGT CTTNCAACAC
301 ANACNCTATC TTGACGACTN CAAAAAANAA CACAATCTGG ACATACCNN
351 AGTCTTNCAA GTGCCGACCG CGCCTTTGGG ACTGTACCCG GGCAAGCTGA
401 AATCGCTGGA NNAAGTCAAA GANGGCAGCA CCGTATCCGC GCCCAACGAC
451 CCGTNNNACT TCGNCCGCGT CTGGTGATG CTCGACGAAC TGGGTTNGAT
50 501 CAAACTCAAA GACNGCATCA NNNNGNNGNN NNNANCNANA NNNGANANN
551 NNNNANNNT NNNNNNNNN NNNNNCNCG NNNNNNANN NNNNNNNNN
601 NCGNNTNNNN NNGCNNNNNT NNANNNTNNN NNCNNCNNNN NNNNTNNNN
651 NANNANNAGC GGCATGAAGC TGACCGAAGC CCTGTTCCAA GAACCGAGCT
701 TTGCCTATGT CAACTGGTCT GCCGTCAAAA CCGCCGACAA AGACAGCCAA
55 751 TGGCTTAAAG ACGTAACCGA GGCCTATAAC TCCGACGCGT TCAAAGCCTA
801 CGCGCACAAA CGCTTCGAGG GCTACAAATC CCCTGCCGCA TGAATGAAG
851 GCGCAGCCAA ATAA

```

This is predicted to encode a protein having amino acid sequence <SEQ ID 220>:

```

1 MKTFFKTLSA AALALILAAAC GGQKDSAPAA SASAAADNGA AXKEIVFGTT

```

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51 VGDFGDMVKE XIQPELEKKG YTVKLVEFTD YVRXNLALAE GELDINVXQH
 101 XXYLDDXKKX HNLDITXVXQ VPTAPLGLYP GKLSLXXVK XGSTVSAPND
 151 PXXFXRVLM LDELGXIKLK DXIXXXXXXX XXXXXXXXXX XXXXXXXXXX
 201 XXXAXXXXXX XXXXXXXXXX GMKLTEALFQ EPSFAYVNW AVKTADKDSQ
 251 WLKDVTEAYN SDAFKAYAHK RFEGYKSPAA WNEGAAK*

A leader peptide is underlined.

Further analysis of these strain A sequences revealed the complete DNA sequence <SEQ ID 221>:

1 ATGAAAACCT TCTTCAAAAC CCTTCCGCC GCCGCACTCG CGCTCATCCT
 51 CGCGCGCTGC GCGGCTCAA AAGATAGCG CCGCGCCGCA TCCGCTCTG
 101 CCGCGCCGCA CAACGGCGCG GCGAAAAAG AAATCGTCTT CGGCACGACC
 151 GTCGGCGACT TCGCGATAT GGTCAAAGAA CAAATCCAAC CCGAGCTGGA
 201 GAAAAAGGC TACACCGTCA AACTGGTCTGA GTTACCGAC TATGTGCGCC
 251 CGAATCTGGC ATTGGCTGAG GCGAGTTGG ACATCAACGT CTTCCAACAC
 301 AAACCCATC TTGACGACTT CAAAAAGAA CACAATCTGG ACATCACCAG
 15 351 AGTCTTCCAA GTGCCGACCG CGCCTTTGGG ACTGTACCGG GGCAAGCTGA
 401 AATCGCTGGA AGAAGTCAA GACGGCAGCA CCGTATCCGC GCCCAACGAC
 451 CCGTCCAAC TCGCCCGCGT CTTGGTGATG CTCGACGAAC TGGGTGGAT
 501 CAAACTCAA GACGGCATCA ATCCGCTGAC CGCATCCAAA GCGGACATTG
 551 CCGAAAACCT GAAAAACATC AAAATCGTCG AGCTTGAAGC CGCGCAACTG
 20 601 CCGCGTAGCC GCGCCGACGT GGATTTTGGC GTCGTCAACG GCAACTACGC
 651 CATAAGCAGC GGCATGAAGC TGACCGAAGC CCTGTTCCAA GAACCGAGCT
 701 TTGCCTATGT CAACTGGTCT GCCGTCAAAA CCGCCGACAA AGACAGCCAA
 751 TGGCTTAAAG ACGTAACCGA GGCCTATAAC TCCGACGCGT TCAAAGCCTA
 801 CGCGCACAAA CGCTTCGAGG GCTACAAATC CCCTGCCGCA TGGAATGAAG
 25 851 GCGCAGCCAA ATAA

This encodes a protein having amino acid sequence <SEQ ID 222; ORF4a-1>:

1 MKTFFKTLSA AALALILAAC GGQKDSAPAA SASAAADNGA AKKEIVFGTT
 51 VGDFGDMVKE QIQPELEKKG YTVKLVEFTD YVRPNLALAE GELDINVXQH
 101 KPYLDDFKKE HNLDITEVFO VPTAPLGLYP GKLSLEEVK DGSTVSAPND
 151 PSNFARVLM LDELGWIKLK DGINPLTASK ADIAENLNKI KIVELEAAQL
 201 PRSRADVDF VVNGNYAISS GMKLTEALFQ EPSFAYVNW AVKTADKDSQ
 251 WLKDVTEAYN SDAFKAYAHK RFEGYKSPAA WNEGAAK*

ORF4a-1 and ORF4-1 show 99.7% identity in 287 aa overlap:

35	orf4a-1	10	20	30	40	50	60
		MKTFFKTLSAALALILAACGGQKDSAPAAASASAAADNGAAKKEIVFGTTVGDFGDMVKE					
	orf4-1	MKTFFKTLSAALALILAACGGQKDSAPAAASASAAADNGAAKKEIVFGTTVGDFGDMVKE					
40	orf4a-1	70	80	90	100	110	120
		QIQPELEKKG YTVKLVEFTD YVRPNLALAE GELDINVXQH KPYLDDFKKE HNLDITEVFO					
	orf4-1	QIQAELEKKG YTVKLVEFTD YVRPNLALAE GELDINVXQH KPYLDDFKKE HNLDITEVFO					
45	orf4a-1	130	140	150	160	170	180
		VPTAPLGLYP GKLSLEEVK DGSTVSAPND PSNFARVLM LDELGWIKLK DGINPLTASK					
	orf4-1	VPTAPLGLYP GKLSLEEVK DGSTVSAPND PSNFARVLM LDELGWIKLK DGINPLTASK					
50	orf4a-1	190	200	210	220	230	240
		ADIAENLNKI KIVELEAAQL PRSRADVDF VVNGNYAISS GMKLTEALFQ EPSFAYVNW					
	orf4-1	ADIAENLNKI KIVELEAAQL PRSRADVDF VVNGNYAISS GMKLTEALFQ EPSFAYVNW					
55	orf4a-1	250	260	270	280		
		AVKTADKDSQ WLKDVTEAYN SDAFKAYAHK RFEGYKSPAA WNEGAAKX					
	orf4-1	AVKTADKDSQ WLKDVTEAYN SDAFKAYAHK RFEGYKSPAA WNEGAAKX					
60		250	260	270	280		

[illegible]

ORF4 shows 93.6% identity over a 94aa overlap with a predicted ORF (ORF4.ng) from *N. gonorrhoeae*:

```

25      orf4nm.pep                                10                20                30
                                                MKTFFKTL$AAALALI$AACGXQKDSAPAA
                                                |||||:::||:|||||
orf4ng    RANAVXTPNPDGRTPLCSFLFETATTSGENMKTFFKTLSTASLALI$AACGGQKDSAPAA
              200            210            220            230            240            250

30          40           50           60           70           80           89
orf4nm.pep SASA-AADNGAAKKEIVFGTTVGDFGDMVKEQIQAELEKKGYTVKLVEFTDYVRPNLALA
         |: :|||||||||||||||||||||
orf4ng     SAAAPSADNGAAKKEIVFGTTVGDFGDMVKEQIQAELEKKGYTVKLVEFTDYVRPNLALA
             260           270           280           290           300           310

35          90
orf4nm.pep EGE$
        |||
orf4ng     EGELDINV$QH$KPYLDDFKKEHNLDITEAFQVP$TAP$GLYPGKLSLEEVKDGSTVSAPN
               320           330           340           350           360           370
```

40 The complete length ORF4ng nucleotide sequence <SEQ ID 223> was predicted to encode a protein having amino acid sequence <SEQ ID 224>:

45

1	MKTFFKTLST	ASLALILAA <u>C</u>	GGQKDSAPAA	SAAAPSADNG	AAKKEIVFGT
51	TVGDFGDMVK	E <u>Q</u> IQAEELEKK	GYTVKLVEFT	PYVRLPNLALA	EGELDINVFGT
101	HKPYLDDFKK	EHNLDLTFEAF	QVPTAPLGLY	DGKLSLEEV	KQGSTVSAPN
151	DPSNFARALV	MLNELGWIKL	KDGINPLTAS	KADIAENLKN	IKIVELEAAQ
201	LPRSRADVDF	AVVNGNYAIS	SGMKLLEALF	QEPSFAYVNW	SAVKTADKDS
251	OWLKDVTEAY	NSDAFKAYAH	KRFEGYKYPA	AWNEGAAK*	

Further analysis revealed the complete length ORF4ng DNA sequence <SEQ ID 225> to be:

50	1	atgAAAACCT	TCTTCAA AAC	cctttccgc	gccgcaCTCG	CGCTCATCCT
	51	CGCAGCCTGc	ggCggtcaAA	AAGACAGCGC	GCCCgaagcc	tctgcCGCCG
	101	CCCCTTCTGc	CGATAACGgc	gCgGCGAAAA	AAGAAAtcgt	ctTCGGCACG
	151	Accgtgggc	acttcggcgA	TatggTCAAA	GACAAATCC	AagcCGAgct
55	201	gGAGAAAAAA	GgctACACcg	tcAAattggt	cgaatttacc	gactagtgtGC
	251	gCCCGAATCT	GGCATTGGCG	GAGGGCGAGT	TGGACATCAA	CGTCTTCCAA
	301	CACAAACCCt	ATCTTGACGA	TTTCAAAAAA	GAACACAACC	TGGACATCAC
	351	CGAAGCCTTC	CAGTGCCGA	CCGCGCCTTT	GGGACTGTAT	CCGGGCCAAC
60	401	TGAAATCGCT	GGAAGAAGTC	AAAGACGGCA	GCACCGTATC	CGCGCCCAac
	451	gACcgTCCA	ACTTCGCGCA	CGCCTTGGTG	ATGCTGAACG	AAC TGGGTTG
	501	GATCAAATC	AAAGACGGCA	TCAATCCGCT	GACCGCATCC	AAAGCCGACA
	551	TCGCGGAAAA	CTGAAAAAAC	ATCAAAATCG	TCGAGCTTGA	ACCGCGCAAA

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5
 601 CTGCCGCGCA GCCGCGCCGA CGTGGATTTT GCCGTCGTCA ACGGCAACTA
 651 CGCCATAAGC AGCGGCATGA AGCTGACCGA AGCCCTGTTC CAAGAGCCGA
 701 GCTTTGCCTA TGTCAACTGG TCTGCCgtcA AAACCGCCGA CAAAGACAGC
 751 CAATGGCTTA AAGACGTAAC CGAGGCCTAT AACTCCGACG CGTTCAAAGC
 801 CTACGCGCAC AAACGCTTCG AGGGCTACAA ATACCTGTCC GCATGGAATG
 851 AAGGCGCAGC CAAATAA

This encodes a protein having amino acid sequence <SEQ ID 226; ORF4ng-1>:

10
 1 MKTFFKTLSA AALALILAAC GGQKDSAPAA SAAAPSDNG AAKKEIVFGT
 51 TVGDFGDMVK EQIQAELEKK GYTVKLVEFT DYVRENLALA EGELDINVFO
 101 HKPYLDDFKK EHNLDITEAF QVPTAPLGLY PGKLKSLLEV KDGSTVSAPN
 151 DPSNFARALV MLNELGWIKL KDGINPLTAS KADIAENLKN IKIVELEAAQ
 201 LPRSRADVDF AVVNGNYAIS SGMKLTEALF QEPSFAYVNW SAVKTADKDS
 251 QWLKDVTEAY NSDAFKAYAH KRFEGYKYP AAWNEGAAK*

This shows 97.6% identity in 288 aa overlap with ORF4-1:

15
 orf4-1.pep 10 20 30 40 50 59
 MKTFFKTLSAALALILAACGGQKDSAPAAASA-AADNGAAKKEIVFGTTVGDFGDMVK
 orf4ng-1 MKTFFKTLSAALALILAACGGQKDSAPAAASAAAPSDNGAAKKEIVFGTTVGDFGDMVK
 20
 orf4-1.pep 60 70 80 90 100 110 119
 EQIQAELEKKGYTVKLVEFTDYVRPNLALAEGELDINVFOHKPYLDDFKKEHNLDITEVF
 orf4ng-1 EQIQAELEKKGYTVKLVEFTDYVRPNLALAEGELDINVFOHKPYLDDFKKEHNLDITEAF
 25
 orf4-1.pep 120 130 140 150 160 170 179
 QVPTAPLGLYPGKLKSLLEVKGSTVSAPNDPSNFARVLVMLDELGWIKLKDGINPLTAS
 orf4ng-1 QVPTAPLGLYPGKLKSLLEVKGSTVSAPNDPSNFARALVMLNELGWIKLKDGINPLTAS
 30
 orf4-1.pep 180 190 200 210 220 230 239
 KADIAENLKNIKIVELEAAQLPRSRADVDFAVVNGNYAISSGMKLTEALFQEPSFAYVNW
 orf4ng-1 KADIAENLKNIKIVELEAAQLPRSRADVDFAVVNGNYAISSGMKLTEALFQEPSFAYVNW
 35
 orf4-1.pep 240 250 260 270 280
 SAVKTADKDSQWLKDVTEAYNSDAFKAYAHKRFEGYKSPAANNEGAAKX
 orf4ng-1 SAVKTADKDSQWLKDVTEAYNSDAFKAYAHKRFEGYKYPAAWNEGAAKX
 40
 orf4-1.pep 250 260 270 280
 SAVKTADKDSQWLKDVTEAYNSDAFKAYAHKRFEGYKSPAANNEGAAKX
 orf4ng-1 SAVKTADKDSQWLKDVTEAYNSDAFKAYAHKRFEGYKYPAAWNEGAAKX

45 In addition, ORF4ng-1 shows significant homology with an outer membrane protein from the database:

50
 ID LIP2_PASHA STANDARD; PRT; 276 AA.
 AC Q08869;
 DT 01-NOV-1995 (REL. 32, CREATED)
 DT 01-NOV-1995 (REL. 32, LAST SEQUENCE UPDATE)
 DT 01-NOV-1995 (REL. 32, LAST ANNOTATION UPDATE)
 DE 28.2 KD OUTER MEMBRANE PROTEIN PRECURSOR. . . .
 SCORES Init1: 279 Initn: 416 Opt: 494
 Smith-Waterman score: 494; 36.0% identity in 275 aa overlap
 55
 orf4ng-1.pep 10 20 30 40 50
 MKTFFKTLSAAL--ALILAACGGQKDSAPAAASAAAPSDNGAAKKEIVFGTTVGDFGDM
 lip2_pasha MNFKKLLGVALVSALALTACKDEKAQAPATTA---KTENKAPLK---VGVMTGPEAQM
 60
 orf4ng-1.pep 60 70 80 90 100 110
 VKEIQAELEKKGYTVKLVEFTDYVRPNLALAEGELDINVFOHKPYLDDFKKEHNLDITE
 :: :: || | |::|::|::| | | :|| |::| |::| ::

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	lip2_pasha	TEVAVKIAKEKYGLDVELVQFTEYTPQPNAAHLSKDLNANAFQTPVYLEQEVKDRGYKLAI	60	70	80	90	100	110
5	orf4ng-1.pep	AFQVPTAPLGLYPGKLSLEEVKDGSTVSAPNDPSNFARALVMLNELGWIKLKDGINPLT	120	130	140	150	160	170
	lip2_pasha	IGNTLVWFPIAAYSKIKNISELKDGATVAIPNNASNTARALLLLOAHGLLKLKDPKN-VF	120	130	140	150	160	170
10	orf4ng-1.pep	ASKADIAENLNKNIKIVELEAAQLPRSRADVDFAVVNGNYAISSGMKLTE--ALFQEPSFA	180	190	200	210	220	230
	lip2_pasha	ATENDIENPKNIKIVQADTSLLTRMLDDVELAVINNTYAGQAGLSPDKDGIIVESKDSP	180	190	200	210	220	230
15	orf4ng-1.pep	YVNWSAVKTADKDSQWLKDVTEAYNSDAFKAYAHKRFEQYKYPAAWNEGAAX	240	250	260	270	280	289
20	lip2_pasha	YVNLVVSREDNKDDPRLQTFVKSFQTEEVFQEALKLFNGGVVKGW	240	250	260	270		

Based on this analysis, including the homology with the outer membrane protein of *Pasteurella haemolittica*, and on the presence of a putative prokaryotic membrane lipoprotein lipid attachment site in the gonococcal protein, it was predicted that these proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

ORF4-1 (30kDa) was cloned in pET and pGex vectors and expressed in *E.coli*, as described above. The products of protein expression and purification were analyzed by SDS-PAGE. Figures 8A and 8B show, respectively, the results of affinity purification of the His-fusion and GST-fusion proteins. Purified His-fusion protein was used to immunise mice, whose sera were used for ELISA (positive result), Western blot (Figure 8C), FACS analysis (Figure 8D), and a bactericidal assay (Figure 8E). These experiments confirm that ORF4-1 is a surface-exposed protein, and that it is a useful immunogen.

Figure 8F shows plots of hydrophilicity, antigenic index, and AMPHI regions for ORF4-1.

Example 27

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 227>:

	1	CCTCGTCGTC	CTCGGCATGC	TCCAGTTTCA	AGGGGCGATT	TACTCCAAGG
	51	CGGTGGAACG	TATGCTCGGC	ACGGTCATCG	GGCTGGGCGC	GGGTTTGGGC
	101	GTTTTATGGC	TGAACAGCA	TTATTTCCAC	GGCAACCTCC	TCTTCTACCT
40	151	CACCGTCGCG	ACGGCAAGCG	CACTGGCCGG	CTGGGCGGCG	GTGGGCAAAA
	201	ACGGCTACGT	CCCTmTGCTG	GCAGGGCTGA	CGATGTGTAT	GCTCATCGGC
	251	GACAACGGCA	GCGAATGGCT	CGACAGCGGA	CTCATGCGCG	CCATGAACGT
	301	CCTCATCGGC	GyGGCCATCG	CCATCGCCGC	CGCCAACTG	CTGCCGCTGA
	351	AATCCACACT	GATGTGGCGT	TTCATGCTTG	CCGACAACCT	GGCCGACTGC
45	401	AGCAAAATGA	TTGCCGAAAT	CAGCAACGGC	AGGCGCATGA	CCCGCGAACG
	451	CCTCGAGGAG	AACATGGCGA	AAATGCGCCA	AATCAACGCA	CGCATGGTCA
	501	AAAGCCGCAG	CCATCTCGCC	GCCACATCGG	GCGAAAGCTG	CATCAGCCCC
	551	GCCATGATGG	AAGCCATGCA	GCACGCCAC	CGTAAATCG	TCAACACCAC
	601	CGAGCTGCTC	CTGACCACCG	CCGCCAAGCT	GCAATCTCCC	AAACTCAACG

5 651 GCAGCGAAAT CCGGCTGCTT GACCGCCACT TCACACTGCT CCAAAC....
 701 GC AGACACGCCC GCCGCATCCG
 751 CATCGACACC GCCATCAACC CCGAACTGGA AGCCCTCGCC GAACACCTCC
 801 ACTACCAATG GCAGGGCTTC CTCTGGCTCA GCACCGATAT GCGTCAGGAA
 851 ATTTCCGCCC TCGTCATCCT GCTGCAACGC ACCCGCCGCA AATGGCTGGA
 901 TGCCCACGAA CGCCAACACC TCGCCAAAG CCTGCTTGA

This corresponds to the amino acid sequence <SEQ ID 228; ORF8>:

10 1PRRP RHAPVSRGDL LQGGGYARH GHRAGRGFGR FMAEPALFPR
 51 QPPLLPHRRH GKRTGRLGGG RQKRLREXAG RADDVYAHRR QRQRMARQRT
 101 HARHERPHRR GHRHRRRQTA AAEIHTDVAF HACRQPGRLQ QNDCRNQQRQ
 151 AHDPRTPRGE HGENAPNORT HGQKPQPSRR HIGRKLHQPR HDGSHAARPP
 201 XNRQHHRAAP DHRROAAISQ TQRQRNPAAX PPLHTAPN...Q
 251 TRPPHPRHRH HQPRTGSPRR TPPLPMAGLP LAQHRYASGN FRPRHPAATH
 301 PPQMAGCPRT PTPAPKPA*

15 Computer analysis of this amino acid sequence gave the following results:

Sequence motifs

ORF8 is proline-rich and has a distribution of proline residues consistent with a surface localization. Furthermore the presence of an RGD motif may indicate a possible role in bacterial adhesion events.

20 Homology with a predicted ORF from *N.gonorrhoeae*

ORF8 shows 86.5% identity over a 312aa overlap with a predicted ORF (ORF8.ng) from *N. gonorrhoeae*:

25 orf8ng 1 MDRDDLRLRRPHAPVPRDL LQGGGYARYGHRAGRGFGRFMAEPALFPR 50
 orf8.pep 1PRRP RHAPVSRGDL LQGGGYARH GHRAGRGFGRFMAEPALFPR 44
 orf8ng 51 QPPLLPHRRHGKRTGRLGGGRQKRLRPYVGADDVHAHRRQRQRMARQRP 100
 orf8.pep 45 QPPLLPHRRHGKRTGRLGGGRQKRLRPXAGRADDVYAHRRQRQRMARQRT 94
 30 orf8ng 101 DARDERPHRRHRHRCRRQTAAAEIHTDVAFHACRQPGRLQ QNDCRNQQRQ 150
 orf8.pep 95 HARHERPHRRGHRHRRRQTA AAEIHTDVAFHACRQPGRMQ QNDCRNQQRQ 144
 35 orf8ng 151 AYDARTFGAEYQONAPNORTHGQKPQPPRRHIGRKPHQPLHDGSHAARPP 200
 orf8.pep 145 AHDPRTPRGEHGENAPNORTHGQKPQPSRRHIGRKLHQPRHDGSHAARPP 194
 40 orf8ng 201 QNRQHHRAAPDHRROAAISQTQRQRNPAARPPLHTAPNRPATNRRPHQRQ 250
 orf8.pep 195 XNRQHHRAAPDHRROAAISQTQRQRNPAAXPPLHTAPN.....Q 244
 orf8ng 251 TRPPHPRHRHQPRTGSPRRTPPLPMAGFPLAQHRYASGNFRPRHPATH 300
 45 orf8.pep 245 TRPPHPRHRHQPRTGSPRRTPPLPMAGLPLAQHRYASGNFRPRHPAATH 294
 orf8ng 301 PPQMAGCPRTPTPAPKPA* 319
 orf8.pep 295 PPQMAGCPRTPTPAPKPA* 313

50 The complete length ORF8ng nucleotide sequence <SEQ ID 229> is predicted to encode a protein having amino acid sequence <SEQ ID 230>:

55 1 MDRDDLRLRR RHAPVPRDL LQGGGYARY GHRAGRGFGR FMAEPALFPR
 51 QPPLLPHRRH GKRTGRLGGG RQKRLRPYVG GADDVHAHRR QRQRMARQRP
 101 DARDERPHRR RHRHRRRQTA AAEIHTDVAF HACRQPGRLQ QNDCRNQQRQ
 151 AYDARTFGAE YQONAPNORT HGQKPQPPRR HIGRKPHQPL HDGSHAARPP


```

201 QNRQHHRAAP DHRROAAISQ TQRQRNPAAR PPLHTAPNRP ATNRRPHQRQ
251 TRPPHPRHRH HQPRTGSPRR TPPLPMAGFP LAQHQYASGN FRPRHPPATH
301 PPQMAGCPRT PTPAPKPA*

```

Based on the sequence motifs in these proteins, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 28

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 231>:

```

10      1 ..GAAATCAGCC TGC GGTC CGA CNACAGGCCG GTTCCGTGN CGAAGCGGCG
      51 GGATTCGGA CTTTTCTGC TGTGGACGG CGGCAACAGC CGGCTCAAGT
      101 GGGCGTGGT GGAAAACGGC ACGTTCGCAA CCGTCGGTAG CGCGCCGTAC
      151 CGCGATTGT CGCCTTTGGG CGCGGAGTGG GCGGAAAAGG CGGATGGAAA
      201 TGTCCGCATC GTCGGTTGCG CTGTGTGCGG AGAATTCAAA AAGGCACAAG
      251 TGCAGGAACA GCTCGCCCGA AAAATCGAGT GGCTGCCGTC TTCCGCACAG
      15 GCTTT.GGCA TACGCAACCA CTACCGCCAC CCCGAAGAAC ACGGTTCCGA
      301 CCGCTGGTTC AACGCCTTGG GCAGCCGCCG CTTAGCCGC AACGCCTGCG
      351 TCGTCGTCAG TTGCGGCACG GCGGTAACGG TTGACGCGCT CACCGATGAC
      401 GGACATTATC TCGGAGA.GG AACCATCATG CCGGTTTCC ACCTGATGAA
      451 AGAATCGCTC GCCGTCCGAA CCGCCAACCT CAACCGGCAC GCCGTAAGC
      501 GTTATCCTTT CCCGACCGG..
      551

```

This corresponds to the amino acid sequence <SEQ ID 232; ORF61>:

```

      1 ..EISLRSDXRP VSVXKRRDSE RFLLLDGGNS RLKWAVVENG TFATVGSAPY
      51 RDLSPGLAEW AEKADGNVRI VGCAVCGEFK KAQVQEQLAR KIEWLPSSAQ
      101 AXGIRNHYRH PEEHGS DRWF NALGSRFRSR NACVVVSCGT AVTVDALTD
      25 GHYLGXGTIM PGFHLMKESL AVRTANLNRH AGKRYPFPT..
      151

```

Further work revealed the complete nucleotide sequence <SEQ ID 233>:

```

      1 ATGACGGTTT TGAAGCTTTC GCACTGGCGG GTGTGGCGG AGCTTGCCGA
      51 CGGTTGCGG CAACACGTCT CGCAACTGGC GCGTATGGCG GATATGAAGC
      101 CGCAGCAGCT CAACGGTTTT TGGCAGCAGA TGCCGGCGCA CATACGCGGG
      30 CTGTTGCGCC AACACGACGG CTATTGGCGG CTGGTGCGCC CATTGGCGGT
      201 TTTGATGCG GAAGTTTTCG GCGAGCTGGG GGAAAGGTCG GGTTTTCAGA
      251 CGGCATTGAA GCACGAGTGC GCGTCCAGCA ACGACGAGAT ACTGGAATTG
      301 GCGCGGATTG CGCCGGACAA GGCGCACAAA ACCATATGCG TGACCCACCT
      351 GCAAAGTAAG GGCAGGGGGC GGCAGGGGGC GAAGTGGTCG CACCGTTTGG
      35 GCGAGTGTCT GATGTTTCAG TTTGGCTGGG TGTTTGACCG GCCGAGTAT
      401 GAGTTGGGTT CGCTGTGCGC TGTTGCGGCA GTGGCGTGTC GGCGCGCCTT
      451 GTCGCGTTTA GGTTTGGATG TGCAGATTAA GTGGCCCAAT GATTTGGTTG
      501 TCGGACGCGA CAAATTGGGC GGCATTCTGA TTGAAACGGT CAGGACGGGC
      551 TCGAAAACGG TTGCCGTGGT CCGTATCGGC ATCAATTTTG TCCTGCCCAA
      601 GGAAGTAGAA AATGCCGCTT CCGTGCAATC GCTGTTTCAG ACGGCATCGC
      40 GCGCGGGCAA TGCCGATGCC GCGGTGCTGC TGGAAACGCT GTTGGTGGAA
      701 CTGGACGCGG TGTGTTGTC AATATGCGCG GACGGATTTG CGCCTTTTGT
      751 GCGGGAATAT CAGGCTGCCA ACCGCGACCA CGGCAAGGCG GTATTGCTGT
      801 TGCGCGACGG CGAAACCGTG TTCGAAGGCA CGGTTAAAGG CGTGGACGGA
      45 CAAGGCGTTT TGCATTGGA AACGCGAGAG GGCAAACAGA CGGTCGTGAG
      901 CGCGGAAATC AGCCTGCGGT CCGACGACAG GCCGTTTTC GTGCCGAAGC
      1001 GCGCGGATTG GGAACGTTTT CTGCTGTTGG ACGGCGGCAA CAGCCGGCTC
      1051 AAGTGGGCGT GGGTGGAAAA CGGCACGTTT GCAACCGTCG GTAGCGCGCC
      1101 GTACCGCGAT TTGTGCGCTT TGGGCGCGGA GTGGGCGGAA AAGGCGGATG
      50 GAAATGTCCG CATCGTGGT TCGCTGTGT GCGGAGAATT CAAAAGGCA
      1151 CAAGTGCAGG AACAGCTCGC CCGAAAAATC GAGTGGCTGC CGTCTTCCGC
      1201 ACAGGCTTTG GGCATACGCA ACCACTACCG CCACCCGAA GAACACGGTT
      1251 CCGACCGCTG GTTCAACGCC TTGGGCGAGC GCCGCTTCAG CCGCAACGCC
      1301 TGCGTCGTCG TCAGTTGCGG CACGGCGGTA ACGGTTGACG CGCTCACCGA
      1351 TGACGGACAT TATCTCGGGG GAACCATCAT GCCCGGTTTC CACCTGATGA
      55 AAGAATCGCT CGCCGTCCGA ACCGCCAACC TCAACCGGCA CGCCGTAAG
      1401 CGTTATCCTT TCCCAGCCAC AACGGGCAAT GCCGTCGCCA GCGGCATGAT
      1451 GGATGCGGTT TGCGGCTCGG TTATGATGAT GCACGGGCGT TTGAAAGAAA
      1501 AAACCGGGGC GGGCAAGCCT GTCGATGTCA TCATTACCGG CGGCGGCGCG
      1601

```

1651 GCAAAAGTTG CCGAAGCCCT GCCGCCTGCA TTTTGGCGG AAAATACCGT
 1701 GCGCGTGGCG GACAACCTCG TCATTACGG GTTGTGAAC ATGATTGCCG
 1751 CCGAAGGCAG GGAATATGAA CATATTAA

This corresponds to the amino acid sequence <SEQ ID 234; ORF61-1>:

5 1 MTVLKLSHWR VLAELADGLP QHVSQ LARMA DMKPQQLNGF WQQMPAHIRG
 51 LLRQHDGYWR LVRPLAVFDA EGLREL GERS GFQTALKHEC ASSNDEILEL
 101 ARIAPDKAHK TICVTHLQSK GRGRQGRKWS HRLGECLMFS FGWVFD RPQY
 151 ELGSLSPVAA VACRRALSRL GLDVQIKWPN DLVVGRDKLG GILIETVRTG
 201 GKTAVAVVGIG INFVLPKEVE NAASVQSLEFQ TASRRGNADA AVLLETLLVE
 10 251 LDAVLLQYAR DGFAPFVAEY QAANRDHGKA VLLLRDGETV FEGTVKGVDG
 301 QGVLHLETAE GKQTVVSGEI SLRSDDRPVS VPKRRDSERF LLLDGGNSRL
 351 KWAUVENGTF ATVGSA PYRD LSPLGAEWAE KADGNVRIVG CAVCGEFKKA
 401 QVQEQ LARKI EWLPSSAQAL GIRNHYRHPE EHGS DRWFNA LGSRRFRSRNA
 451 C VVVSCGTAV TVDALTD DGH YLGGT IMPGF HLMKESLAVR TANLNRHAGK
 15 501 RYFPPTTGN AVASGMMDAV CGSVMMHGR LKEKTGAGKP VDVIIITGGGA
 551 AKVAEALPPA FLAENTVRVA DNLVIYGLLN MIAAEGREYE HI*

Figure 9 shows plots of hydrophilicity, antigenic index, and AMPHI regions for ORF61-1. Further computer analysis of this amino acid sequence gave the following results:

Homology with the baf protein of *B. pertussis* (accession number U12020).

20 ORF61 and baf protein show 33% aa identity in 166aa overlap:

orf61 23 LLLDGGNSRLKWAWE-NGTFATVGSA PYR----DLSP LGAEWAEKADGNVRIVGCAVCG 77
 +L+D GNSRLK W + + A AP DL LG A R +G V G
 baf 3 ILIDSGNSRLKVGWFD PDAPQAAREPAPVAFDNL DLDALGRWLATLPRRQ RALGVNVAG 62
 25 orf61 78 EFKKAQVQEQ LAR---KIEWLPSSAQAXGIRNHYRHPEEHGSDRW---FNALGSRFRSRN 131
 + + L I WL + A G+RN YR+P++ G+DRW L +
 baf 63 LARGEAIAATLRAGGCDIRW LRAQPLAMGLRNGYRNP DQLGADRWACMVGV LARQPSVHP 122
 orf61 132 ACVVVSCGTAVTVDALTD DGHYLGXGTIMPGF HLMKESLAVRTANL 177
 +V S GTA T+D + D + G G I+PG +M+ +LA TA+L
 30 baf 123 PLLVASFGTATTLDTIGPDNVFP G-LILPGPAMMRGALAYGTAHL 167

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF61 shows 97.4% identity over a 189aa overlap with an ORF (ORF61a) from strain A of *N.*

35 *meningitidis*:

orf61.pep 10 20 30
 EISLRSDXRPVSVXKRRDSERFLLLDGGNS
 orf61a 290 300 310 320 330 340
 TVFEGTVKGVDGQGV L HLETAEGKQTVVSGEISLRSDDRPVSVPKRRDSERFLLLDGGNS
 40 orf61.pep 40 50 60 70 80 90
 RLKWAUVENGTFATVGSA PYRDLSP LGAEWAEKADGNVRIVGCAVCGEFKKAQVQEQ LAR
 orf61a 350 360 370 380 390 400
 RLKWAUVENGTFATVGSA PYRDLSP LGAEWAEKV DGNVRIVGCAVCGEFKKAQVQEQ LAR
 45 orf61.pep 100 110 120 130 140 150
 KIEWLPSSAQAXGIRNHYRHPEEHGSDRWFNALGSRFRSRNACVVVSCGTAVTVDALTD D
 orf61a 410 420 430 440 450 460
 KIEWLPSSAQALGIRNHYRHPEEHGSDRWFNALGSRFRSRNACVVVSCGTAVTVDALTD D
 50 orf61.pep 160 170 180 189
 GHYLGXGTIMPGF HLMKESLAVRTANLNRHAGKRYPFPT
 orf61a 470 480 490 500 510 520
 GHYLG-GTIMPGF HLMKESLAVRTANLNRHAGKRYPFPTTTGN AVASGMMDAVCGSVMM
 55 orf61a HGR LKEKTGAGKPV DVIITGGGA AKVAEALPPA FLAENTVRVADNLVIHGLLN LIAAEGG
 60

530 540 550 560 570 580

The complete length ORF61a nucleotide sequence <SEQ ID 235> is:

```

1 ATGACGGTTT TGAAGCCTTC GCACTGGCGG GTGTTGGCGG AGCTTGCCGA
5 51 CGGTTTGCCG CAACACGTCT CGCAACTGGC GCGTATGGCG GATATGAAGC
101 CGCAGCAGCT CAACGGTTTT TGGCAGCAGA TGCCGCGCGA CATACGCGGG
151 CTGTTGCGCC AACACGACGG CTATTGGCGG CTGGTGCGCC CATTGGCGGT
201 TTTCGATGCC GAAGGTTTGC GCGAGCTGGG GGAAAGGTCG GGTTTTCAGA
251 CGGCATTGAA GCACGAGTGC GCGTCCAGCA ACGACGAGAT ACTGGAATTG
301 GCGCGGATTG CGCCGGACAA GGCGCACAAA ACCATATGTG TGACCCACCT
10 351 GCAAAGTAAG GGCAGGGGGC GGCAGGGGCG GAAGTGGTCG CACCGTTTGG
401 GCGAGTGTCT GATGTTGAGT TTTGGCTGGG TGTTTGACCG GCCGCAGTAT
451 GAGTTGGGTT CGCTGTGCGC TGTTGCGGCA GTGGCGTGCC GGCGCGCCTT
501 GTCGCGTTTG GGTTTGAAAA CGCAAATCAA GTGGCCAAAC GATTTGGTCG
551 TCGGACGCGA CAAATTGGGC GGCATTCTGA TTGAAACGGT CAGGACGGGC
15 601 GCGAAAACGG TTGCGTGGT CCGTATCGCG ATCAATTTCT TGCTGCCCAA
651 GGAAGTGGAA AACGCGCGTT CCGTGCAATC GCTGTTTCAG ACGGCATCGC
701 GCGCGGGAAA TGCCGATGCC GCCGTGTTGC TGGAACGCTT GTTGCGGGAA
751 CTTGATGCGG TGTTGTTGCA ATATGCGCGG GACGGATTTG CGCCTTTTGT
801 GCGCGAATAT CAGGCTGCCA ACCGCGACCA CGGCAAGGCG GTATTGCTGT
20 851 TGCGCGACGG CGAAACCGTG TTCGAAGGCA CGGTTAAAGG CGTGGACGGA
901 CAAGGCGTTC TGCACTTGGA AACGCGCAGG GGCAAACAGA CGGTCGTGAG
951 CGGCGAAATC AGCCTGCGGT CCGACGACAG GCCGGTTTCC GTGCCGAAGC
1001 GCGCGGATTG GGAACGTTTT CTGCTGTTGG ACGGCGGCAA CAGCCGGCTC
1051 AAGTGGGCGT GGGTGGAAAA CGGCACGTTT GCAACCGTCG GTAGCGCGCC
25 1101 GTACCGCGAT TTGTCGCGCT TGCGCGCGGA GTGGGCGGAA AAGGTGGATG
1151 GAAATGTCCG CATCGTCGGT TGCGCGGTGT GCGGAGAATT CAAAAGGCA
1201 CAAGTGCAGG AACAGCTCGC CCGAAAAATC GAGTGGCTGC CGTCTTCCGC
1251 ACAGGCTTTG GGCATACGCA ACCACTACCG CCACCCGAA GAACACGGTT
1301 CCGACCGCTG GTTCAACGCC TTGGGCGAGC GCCGCTTCAG CCGCAACGCC
30 1351 TGCGTCGTCG TCAGTTGCGG CACGCGGGA ACGGTTGACG CGCTCACCGA
1401 TGACGGACAT TATCTCGGGG GAACCATCAT GCCCGGTTTC CACCTGATGA
1451 AAGAATCGCT CGCCGTCCGA ACCGCCAACC TCAACCGGCA CGCCGTAAG
1501 CGTTATCCTT TCCCGACCAC AACGGGCAAT GCCGTCGCA GCGGCATGAT
35 1551 GAGTGCAGGT TGCGGCTCGG TTATGATGAT GCACGGGCGT TTGAAAGAAA
1601 AAACCGGGGC GGGCAAGCCT GTCGATGTCA TCATTACCGG CGGCGGCGCG
1651 GCAAAAGTTG CCGAAGCCCT GCCGCCTGCA TTTTGGCGG AAAATACCGT
1701 GCGCGTGGCG GACAACCTCG TCATTACCGG GCTGCTGAAC CTGATTGCGG
1751 CCGAAGGCGG GGAATCGGAA CATACTTAA

```

This encodes a protein having amino acid sequence <SEQ ID 236>:

```

40 1 MTVLKPSHWR VLAELADGLP QHVSQIARMA DMKPQQLNGF WQMPAHIRG
51 LLRQHDGYWR LVRPLAVFDA EGLRELTERS GFQTALKHEC ASSNDEILEL
101 ARIAPDKAHK TICVTHLQSK GRGRQGRKWS HRLGECIMFS FGWVDFRPQY
151 ELGSLSPVAA VACRRALSRL GLKTQIKWPN DLVVGRDKLG GILLETVRTG
45 201 GKTAVVVGIG INEVLPEKEV NAASVQSLFQ TASRRGNADA AVLLETLLAE
251 LDAVLLQYAR DGFAPFVAEY QAANRDHGKA VLLLRDGETV FEETVKGVVDG
301 QGVLEHLEAE GKQTVVSGEI SLRSDDRPVS VPKRRDSERF LLLDGGNSRL
351 KWAVVENGTG ATVGSAFYRD LSPLGAEWAE KVDGNVRIWG CAVCGEEFKA
401 QVQEQIARKI EWLPSAQAL GIRNHYRHPE EHGSDFWENA LGSRRFSRNA
451 CVVVSCTAV TVDALTDGHH YLGGTIMPGF HLMKESLAVR TANLNRHAGK
50 501 RYFPFTTGN AVASGMMDAV CGSVMMHGR LKEKTGAGKP VDVIITGGGA
551 AKVAEALPPA FLAENTVRVA DNLVIHGLLN LIAAEGGESE HT*

```

ORF61a and ORF61-1 show 98.5% identity in 591 aa overlap:

```

55 orf61a.pep 10 20 30 40 50 60
MTVLKPSHWRVLAELADGLPQHVSQIARMADMKPQQLNGFWQMPAHIRGLLRQHDGYWR
||||| ||||||| ||||||| ||||||| ||||||| |||||||
orf61-1 MTVLKLSHWRVLAELADGLPQHVSQIARMADMKPQQLNGFWQMPAHIRGLLRQHDGYWR
10 20 30 40 50 60

60 orf61a.pep 70 80 90 100 110 120
LVRPLAVFDAEGLRELTERSFGFQTALKHECASSNDEILELARIAPDKAHKTICVTHLQSK
||||| ||||||| ||||||| ||||||| ||||||| |||||||
orf61-1 LVRPLAVFDAEGLRELTERSFGFQTALKHECASSNDEILELARIAPDKAHKTICVTHLQSK
70 80 90 100 110 120

65 130 140 150 160 170 180

```

-174-

5	orf61a.pep	GRGRQGRKWSHRLGECLMFSFGWVFDPRQYELGSLSPVAAVACRRALSRLGLKTQIKWPN	
	orf61-1	GRGRQGRKWSHRLGECLMFSFGWVFDPRQYELGSLSPVAAVACRRALSRLGLDVQIKWPN	
		130 140 150 160 170 180	
10	orf61a.pep	DLVVGRDKLGGILIIETVRTGGKTVAVVGIGINFLPKVEVENAASVQSLFQTASRRGNADA	
	orf61-1	DLVVGRDKLGGILIIETVRTGGKTVAVVGIGINFLPKVEVENAASVQSLFQTASRRGNADA	
		190 200 210 220 230 240	
15	orf61a.pep	AVLLETLLAELDAVLLQYARDGFAPFVAEYQAANRDHGKAVLLLRDGETVFEGTVKGVDG	
	orf61-1	AVLLETLLVELDAVLLQYARDGFAPFVAEYQAANRDHGKAVLLLRDGETVFEGTVKGVDG	
		250 260 270 280 290 300	
20	orf61a.pep	QGVHLHLETAEGKQTVVSSEISLRSDRPVSVPKRRDSEFLLLDGGNSRLKWAWVNGTF	
	orf61-1	QGVHLHLETAEGKQTVVSSEISLRSDRPVSVPKRRDSEFLLLDGGNSRLKWAWVNGTF	
		310 320 330 340 350 360	
25	orf61a.pep	ATVGSAPYRDLSPGLAEWAEEKVDGNVRIVGCAVCGEFKKAQVQEQQLARKIEWLPSSAQAL	
	orf61-1	ATVGSAPYRDLSPGLAEWAEEKADGNVRIVGCAVCGEFKKAQVQEQQLARKIEWLPSSAQAL	
		370 380 390 400 410 420	
30	orf61a.pep	GIRNHRYHPPEEHGSDRWFNALGSRRFSRNACVVVSCGTAVTVDALTDGHHYLGGTIMPGF	
	orf61-1	GIRNHRYHPPEEHGSDRWFNALGSRRFSRNACVVVSCGTAVTVDALTDGHHYLGGTIMPGF	
		430 440 450 460 470 480	
35	orf61a.pep	HLMKESLAVRTANLNRHAGKRYPFPTTTGNAVASGMMDAVCGSVMMHGRLEKKTGAGKP	
	orf61-1	HLMKESLAVRTANLNRHAGKRYPFPTTTGNAVASGMMDAVCGSVMMHGRLEKKTGAGKP	
		490 500 510 520 530 540	
40	orf61a.pep	VDVIITGGGAAKVAEALPPAFLAENTVRVADNLVIHGLLNLIAAEGGESEHTX	
	orf61-1	VDVIITGGGAAKVAEALPPAFLAENTVRVADNLVIYGLLNLMIAAEGREYEHIX	
		550 560 570 580 590	

Homology with a predicted ORF from *N.gonorrhoeae*

ORF61 shows 94.2% identity over a 189aa overlap with a predicted ORF (ORF61.ng) from *N.*

50 *gonorrhoeae*:

55	orf61.pep	EISLRSDXRPVSVXKRRDSEFLLLDGGNS	30
	orf61ng	TVCEGTVKGVDGRGVHLHLETAEGEQTVVSSEISLRPDNRSVSVPKRPDSEFLLLEGGNS	211
60	orf61.pep	RLKWAWVNGTFATVGSAPYRDLSPGLAEWAEEKADGNVRIVGCAVCGEFKKAQVQEQQLAR	90
	orf61ng	RLKWAWVNGTFATVGSAPYRDLSPGLAEWAEEKADGNVRIVGCAVCGESKKAQVKEQLAR	271
65	orf61.pep	KIEWLPSSAQAXGIRNHRYHPPEEHGSDRWFNALGSRRFSRNACVVVSCGTAVTVDALTD	150
	orf61ng	KIEWLPSSAQALGIRNHRYHPPEEHGSDRWFNALGSRRFSRNACVVVSCGTAVTVDALTD	331
	orf61.pep	GHYLGXGTIMPGFHLMKESLAVRTANLNRHAGKRYPFPT	189
	orf61ng	GHYLG-GTIMPGFHLMKESLAVRTANLNRPAKRYPFPTTTGNAVASGMMDAVCGSIMM	390

An ORF61ng nucleotide sequence <SEQ ID 237> was predicted to encode a protein having amino acid sequence <SEQ ID 238>:

```

      1  MFSFGWAFDR PQYELGSLSP VAALACRRAL GCLGLETQIK WPNDLVVGRD
      51  KLGGILITV RAGGKTVAVV GIGINFVLPK EVENAASVQS LFQTASRRGN
1001  ADAAVLLETL LAELGAVLEQ YAEEGFAPFL NEYETANRDH GKAVLLLRDG
151  ETVCEGTVKG VDGRGVLHLE TAEGEQTVVS GEISLRPDNR SVSVPKRPDS
201  ERFLLLLEGGN SRLKWAVVEN GTFATVGSAP YRDLSPLGAE WAEKADGNVR
251  IVGCAVCGES KKAQVKEQLA RKIEWLPSSA QALGIRNHYR HPEEHGSDRW
301  FNALGSRRES RNACVVVSCG TAVTVDALTD DGHYLGGTIM PGFHLMKESL
10    351  AVRTANLNRP AGKRYPFPTT TGNVAVSGMM DAVCGSIMMM HGRLKEKNGA
401  GKPVDVIITG GGAAKVAEAL PPAFLAENTV RVADNLVIHG LLNLIAAEGG
451  ESEHA*

```

Further analysis revealed the complete gonococcal DNA sequence <SEQ ID 239> to be:

```

      1  ATGACGGTTT TGAAGCCTTC GCATTGGCGG GTGTTGGCGG AGCTTGCCGA
15    51  CGGTTTGCCG CAACACGTAT CGCAATTGGC GCGTGAGGCG GACATGAAGC
101  CGCAGCAGCT CAACCGGTTT TGGCAGCAGA TGCCGCGCGA TATACGCGGG
151  CTGTTGCGCC AACACGACGG CTATTGGCGG CTGGTGCGCC CTTGGCGGGT
201  TTTCGATGCC GAAGGTTTGC GCGATCTGGG GGAAAGGTGC GGTTTTCAGA
251  GCGCATTGAA GCACGAGTGC GCGTCCAGCA ACGACGAGAT ACTGGAATTG
20    301  GCGCGGATTG CGCCGGACAA GGCGCACAAA ACCATATGCG TGACCCACCT
351  GCAAAGTAAG GGCAGGGGGC GGCAGGGGGC GAAGTGTCTG CACCGTTTGG
401  GCGAGTGCCT GATGTTCACT TTCGGCTGGG CGTTTGACCG GCCGCAGTAT
451  GAGTTGGGTT CGCTGTCGCC TGTGCGGCA CTGCGTGCC GCGCGCGTTT
501  GGGGTGTTTG GGTGTTGAAA CGCAAATCAA GTGGCCAAAC GATTTGGTCG
25    551  TCGGACCGCA CAAATTGGGC GGCATTCTGA TTGAAACAGT CAGGGCGGGC
601  GGTAACACGG TTGCCGTGGT CCGTATCGGC ATCAATTTCG TGCTGCCCAA
651  GGAAGTGGAA AACGCCGCTT CCGTGCACTC GCTGTTTCAG ACGGCATCGC
701  GGCGGGGCAA TGCCGATGCC GCCGTATTGC TGGAAACATT GCTTGCGGAA
751  CTGGGCGCGG TGTGGAACA ATATGCGGAA GAAGGGTTCG CGCCATTTTT
30    801  AAATGAGTAT GAAACGGCCA ACCGCGACCA CGGCAAGGCG GTATTGCTGT
851  TCGCGCAGCG CGAAACCGTG TCGCAAGGCA CGGTTAAAGC CGTGGACGTA
901  CGAGGCGTTC TGCACTTGGA AACGGCAgaa ggcgaACAGa cggtcgtcag
951  cggcgaaaTC AGcctGCggc ccgacaacAG GTCGGTttcc gtgcccgaagc
1001 gggccggatTC GgaacgtTTT tTGCTgttgg aaggcgggaa cagccgGCTC
35    1051 AAGTGGGCGT GggtggAAAA cggcacgttc gcaaccgtgg gcagcgcgCc
1101 gtaCCGCGAT TTGTCGCTT TGGGCGCGGA GTGGGCGGAA AAGGCGGATG
1151 GAAATGTCCG CATCGTCGGT TGCGCCGTGT GCGGAGAATC CAAAAAGGCA
1201 CAAGTGAAGG AACAGCTCGC CCGAAAAATC GAGTGGCTGC CGTCTTCGCG
1251 ACAGGCTTTC GGCATACGCA ACCACTACCG CCACCCCGAA GAACACGGTT
40    1301 CCGACCGTTG GTTCAACGCC TTGGGCAGCC GCCGCTTCAG CCGCAACGCC
1351 TCGCTCGTCG TCAGTTGCGG CACGGCGGTA ACGGTTGACG CGCTCACCGA
1401 TGACGGACAT TATCTCGGCG GAACCATCAT GCCCGGCTTC CACCTGATGA
1451 AAGAATCGCT CGCCGTCGGA ACCGCCAACC TCAACCGCCC CGCCGGCAAA
45    1501 CGTTACCCTT TCCCGACCAC AACGGGCAAC GCCGTCGCAA GCGGCATGAT
1551 GGACGCGGTT TGCGGCTCGA TAATGATGAT GCACGGCCGT TTGAAAGAAA
1601 AAAACGGCGC GGGCAAGCCT GTCGATGTCA TCATTACCGG CGGCGGCGCG
1651 GCGAAAGTCG CCGAAGCCCT GCCGCCTGCA TTTTGGCGG AAAATACCGT
1701 GCGCGTGGCG GACAACCTCG TCATCCACGG GCTGCTGAAC CTGATTGCCG
1751 CCGAAGGCGG GGAATCGGAA CACGCTTAA

```

50 This corresponds to the amino acid sequence <SEQ ID 240; ORF61ng-1>:

```

      1  MTVLKPSHWR VLAELADGLP QHVSQALAREA DMKPQQLNGF WQMPAHIRG
      51  LLRQHDGYWR LVRPLAVFDA EGLRDLGERS GFQTALKHEC ASSNDEILEL
101  ARIAPDKAHK TICVTHLQSK GRGRQGRKWS HRLGECLMFS FGWAFDRPQY
55    151  ELGSLSPVAA LACRRALGCL GLETQIKWPN DLVVGRDKLG GILITVVRAG
201  GKTAVAVVIG INFVLPKEVE NAASVQSLFQ TASRRGNADA AVLLETLLAE
251  LGAVLEQYAE EGFAPFLNEY ETANRDHGKA VLLLRDGETV CEGTVKGVVDG
301  RGVHLHLETAE GEQTVVSGEI SLRPDNRVS SVKRPDSERF LLEGGNSRL
351  KWAVVENGTF ATVGSAPYRD LSPLGAEWAE KADGNVRIVG CAVCGESKKA
401  QVKEQLARKI EWLPSSAQAL GIRNHRYHPE EHGSDFWNA LGSRRFSRNA
60    451  CVVVSCGTAV TVDALTDGHH YLGGTTPMPG HLMKESLAVR TANLNRPAGK
501  RYFPFTTGN AVASGMMDAV CGSIMMMHGR LKEKNGAGKP VDVIITGGGA
551  AKVAEALPPA FLAENTVRVA DNLVIHGLLN LIAAEGGESE HA*

```

ORF61ng-1 and ORF61-1 show 93.9% identity in 591 aa overlap:

	orf61ng-1.pep	MTVLKPSHWRVLAELADGLPQHVSQALAREADMKPQQLNGFWQQMPAHIRGLLRQHDGYWR	60
	orf61-1	MTVLKLSHWRVLAELADGLPQHVSQALARMADMKPQQLNGFWQQMPAHIRGLLRQHDGYWR	60
5	orf61ng-1.pep	LVRPLAVFDAEGLRDLGERSGFQTALKHECASSNDEILELARIAPDKAHKTICVTHLQSK	120
	orf61-1	LVRPLAVFDAEGLRELGERSGFQTALKHECASSNDEILELARIAPDKAHKTICVTHLQSK	120
10	orf61ng-1.pep	GRGRQGRKWSHRLGECLMFSFGWAFDRPQYELGSLSPVAALACRRALGCLGETQIKWPN	180
	orf61-1	GRGRQGRKWSHRLGECLMFSFGWVDFRPQYELGSLSPVAACRRALSRLGLDVQIKWPN	180
	orf61ng-1.pep	DLVVGRDKLGGILIEITVRAGGKTAVAVVGIGINFVLPKEVENAASVQSLFQTASRRGNADA	240
15	orf61-1	DLVVGRDKLGGILIEITVRTGGKTAVAVVGIGINFVLPKEVENAASVQSLFQTASRRGNADA	240
	orf61ng-1.pep	AVLLETLLAELGAVLEQYAEFGFAPFLNEYETANRDHGKAVLLLRDGETVCEGTVKGVVDG	300
20	orf61-1	AVLLETLLVELDAVLLQYARDGFAPFVAEYQAANRDHGKAVLLLRDGETVFEETVKGVVDG	300
	orf61ng-1.pep	RGVLHLETAEGEQTVVSGEISLRPDNRSVSVPKRPDSEFLLLEGGNSRLKWAVVENGTG	360
	orf61-1	QGVVLHLETAEGKQTVVSGEISLRSDRPVSVPKRRDSEFLLLDGGNSRLKWAVVENGTG	360
25	orf61ng-1.pep	ATVGSAPYRDLSPLGAEWAEEKADGNVRIVGCAVCGESKKAQVKEQLARKIEWLPSSAQAL	420
	orf61-1	ATVGSAPYRDLSPLGAEWAEEKADGNVRIVGCAVCGEFKKAQVQEQQLARKIEWLPSSAQAL	420
30	orf61ng-1.pep	GIRNHYRHPEEHGSDRWFNALGSRRFSRNACVVVSCGTAVTVDALTDGHHYLGTTIMPGF	480
	orf61-1	GIRNHYRHPEEHGSDRWFNALGSRRFSRNACVVVSCGTAVTVDALTDGHHYLGTTIMPGF	480
	orf61ng-1.pep	HLMKESLAVRTANLNRPAKRYFPFPTTGNVAVSGMMDAVCGSIMMMHGRLKEKNAGKGP	540
35	orf61-1	HLMKESLAVRTANLNRHAGKRYFPFPTTGNVAVSGMMDAVCGSVMMMHGRLKEKTGAGKGP	540
	orf61ng-1.pep	VDVIITGGGAAKVAEALPPAFLAENTVRVADNLVIHGLLNLIAAEGGESEHAX	593
40	orf61-1	VDVIITGGGAAKVAEALPPAFLAENTVRVADNLVIYGLLNMIAAEGREYEHIX	593

Based on this analysis, including the homology with the baf protein of *B.pertussis* and the presence of a putative prokaryotic membrane lipoprotein lipid attachment site, it is predicted that these proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

45 Example 29

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 241>:

	1	ATGTTTACC	AAATCCTTGC	CCTGATTATC	TGGAGCAGCT	CGTTTATTGC
	51	CGCCAAATAT	GTCTATGGCG	GCATCGATCC	CGCATTGATG	GTCGGCGTGC
50	101	GCCTGCTAAT	TGCCGCGCTG	CCTGCACTGC	CCGCGTGCCG	CCGTGATGTC
	151	GGCAAGATTC	CGCGTGAGGA	ATGGAAGCCG	TTGCTGATTG	TGTCGTTCCGT
	201	CAACTATGTG	CTGACCTGCG	TGCTTCAGTT	TGTCGGGTTG	AAATACACTT
	251	CCGCCGCCAG	CGCATCGGTC	ATTGTCGGAC	TCGAGCCGCT	GCTGATGGTG
	301	TTTGTCGGAC	ACTTTTCTTT	CAACGACAAA	GCGCGTGCCG	ACCACTGGAT
55	351	ATGCGGCGCG	GCGGCATTTC	CCGGTGTCGC	GCTGCTGATG	GCGGGCGGTG
	401	CGGaAGAGGG	CGGCgaAGTC	GGCTGGTTCG	GCTGCCTGCT	GGTGTGTTTG
	451	GCGGGCGCGG	GCTTTTGTGC	CGCTATGCGT	CCGACGCAAA	GGCTGATTGC
	501	ACGCATCGGC	GCACCGGCAT	TCACATCTGT	TTCCATTGCC	GCCGCATCGT
	551	TGATGTGCCT	GCCGTTTTCG	CTTGCTTTGG	CGCAAAGTTA	TACCGTGGAC
60	601	TGGAGCGTCG	GGATGCTATT	GTCGCTGCTG	TATTGGGTTT	TGGGGTGC..

This corresponds to the amino acid sequence <SEQ ID 242; ORF62>:

-177-

1 MFYQILALII WSSSFIAAKY VYGGIDPALM VGVRLIIAAL PALPACRRHV
 51 GKIPREEWKP LLIVSFVNYV LTLLLQFVGL KYTSAASASV IVGLEPLLMV
 101 FVGHHFFNDK ARAYHWICGA AAFAGVALLM AGGAEEGGEV GWFGCLLVLL
 151 AGAGFCAAMR PTQRILIARIG APAFTSVSIA AASLMCLPFS LALAQSYTVD
 201 WSVGMVLSLL YLGLGC..

Further work revealed the complete nucleotide sequence <SEQ ID 243>:

1 ATGTTTACC AAATCCTGCG CCTGATTATC TGGAGCAGCT CGTTTATGCG
 51 CGCCAAATAT GTCTATGGCG GCATCGATCC CGCATTGATG GTCGGCGTGC
 101 GCCTGCTAAT TGCCGCGCTG CCTGCACTGC CGCCTGCGG CCGTCATGTC
 151 GGCAAGATTC CGCGTGAGGA ATGGAAGCCG TTGCTGATTG TGTCGTTGCT
 201 CAACTATGTG CTGACCTGCG TGCTTCAGTT TGTCGGGTTG AAATACACTT
 251 CCGCCGCCAG CGCATCGGTC ATTGTGCGAC TCGAGCCGCT GCTGATGGTG
 301 TTTGTCGAC ACTTTTTCTT CAACGACAAA GCGCGTGCCT ACCACTGGAT
 351 ATGCGGCGCG GCGGCATTTC CCGGTGTGCG GCTGCTGATG GCGGGCGGTG
 401 CGGAAGAGGG CGGCGAAGTC GGCTGGTTCG GCTGCCTGCT GGTGTTGTTG
 451 GCGGGCGCGG GCTTTTGTGC CGCTATGCGT CCGACGCAA GGCTGATTGC
 501 ACGCATCGGC GCACCGGCAT TCACATCTGT TTCCATTGCC GCGCATCGT
 551 TGATGTGCCT GCCGTTTTTCG CTTGCTTTGG CGCAAAGTTA TACCGTGGAC
 601 TGGAGCGTCG GGATGGTATT GTGCTGCTG TATTTGGGTT TGGGGTGGCG
 651 CTGGTACGCC TATTGGCTGT GGAACAAGGG GATGAGCCGT GTTCTGCCA
 701 ATGTTTCGGG ACTGTTGATT TCGCTCGAAC CCGTCGTCGG CGTGCTGCTG
 751 GCGGTTTTGA TTTTGGGCGA ACACCTGTGC CCCGTGTCCG CTTTGGGCGT
 801 GTTTGTGCTC ATCGCGCCA CTTGGTTGCG CGGCCGGCTG TCGCATCAA
 851 AATAA

25 This corresponds to the amino acid sequence <SEQ ID 244; ORF62-1>:

1 MFYQILALII WSSSFIAAKY VYGGIDPALM VGVRLIIAAL PALPACRRHV
 51 GKIPREEWKP LLIVSFVNYV LTLLLQFVGL KYTSAASASV IVGLEPLLMV
 101 FVGHHFFNDK ARAYHWICGA AAFAGVALLM AGGAEEGGEV GWFGCLLVLL
 151 AGAGFCAAMR PTQRILIARIG APAFTSVSIA AASLMCLPFS LALAQSYTVD
 201 WSVGMVLSLL YLGLGCWYA YWLWNKMSR VPANVSGLLI SLEPVVGVL
 251 AVLILGEHLS PVSALGVFVV IAATLVAGRL SHQK*

Computer analysis of this amino acid sequence gave the following results:

Homology with hypothetical transmembrane protein HI0976 of *H. influenzae* (accession number Q57147)

ORF62 and HI0976 show 50% aa identity in 114aa overlap:

35 Orf62 1 MFYQILALIIWSSSFIAAKYVYGGIDPALMVGVRLIIAALPALPACRRHV
 M YQILAL+IWSSS I K Y +DP L+V VR R KI + K
 HI0976 1 MLYQILALLIWSSSLIVGKLTYSMMDPVLVVQVRLIIAMIIVMPLFLRRWKKIDKPMRKQ 60
 Orf62 61 LLIVSFVNYVLTLLLQFVGLKYTSAASASVIVGLEPLLMVFGVGHFFNDKARAY 114
 L ++F NY LLQF+GLKYTSA+SA ++GLEPLL+VFGVGHFF K +
 40 HI0976 61 LWWLAFENYTAVFLQLQFVGLKYTSAASAVTMIGLEPLLIVFGVGHFFKTKQNGF 114

Homology with a predicted ORF from *N. meningitidis* (strain A)

ORF62 shows 99.5% identity over a 216aa overlap with an ORF (ORF62a) from strain A of *N.*

45 *meningitidis*:

10 20 30 40 50 60
 orf62.pep MFYQILALIIWSSSFIAAKYVYGGIDPALMVGVRLIIAALPALPACRRHV
 50 orf62a MFYQILALIIWSSSFIAAKYVYGGIDPALMVGVRLIIAALPALPACRRHV
 10 20 30 40 50 60
 orf62.pep LLIVSFVNYVLTLLLQFVGLKYTSAASASVIVGLEPLLMVFGVGHFFNDKARAYHWICGA
 55 orf62a LLIVSFVNYVLTLLLQFVGLKYTSAASASVIVGLEPLLMVFGVGHFFNDKARAYHWICGA
 70 80 90 100 110 120
 orf62.pep AAFAGVALLMAGGAEEGGEVWGFCCLLVLLAGAGFCAAMRPTQRILIARIGAPAFTSVSIA
 130 140 150 160 170 180

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```

      |||
orf62a  AAFAGVALLMAGGAEEGGEVGVFGCLLVLLAGAGFCAAMRPTQRLIARIGAPAFTSVSIA
      130      140      150      160      170      180

5      190      200      210
orf62.pep AASLMCLPFSLALAQSYTVDWSVGMVLSLLYLGLGC
      |||
orf62a  AASLMCLPFSLALAQSYTVDWSVGMVLSLLYLGVGCSWYAYWLWNKGMSRVPANVSGLLI
      190      200      210      220      230      240

10     orf62a  SLEPVVGVLAVLILGEHLSPVSVLGVFVVIATLVAGRLSHQKX
      250      260      270      280

```

The complete length ORF62a nucleotide sequence <SEQ ID 245> is:

```

1      ATGTTTACC AAATCCTTGC CCTGATTATC TGGAGCAGCT CGTTTATTGC
15     51      CGCCAAATAT GTCTATGGCG GCATCGATCC CGCATTGATG GTCGGCGTGC
      101     GCCTGCTGAT TGCTGCGCTG CCTGCACTGC CCGCTGCCG CCGTCATGTC
      151     GGCAAGATTC CGCGTGAGGA ATGGAAGCCG TTGCTGATTG TGTGCTTCGT
      201     CAACTATGTG CTGACCCTGC TACTTCAGTT TGTGCGGTG AAATACACTT
20     251     CCGCGGCCAG CGCATCGGTC ATTGTCGGAC TCGAGCCACT GCTGATGGTG
      301     TTTGTCGGAC ACTTTTCTT CAACGACAAA GCGCGTGCCT ACCACTGGAT
      351     ATGCGGCGCG GCGGCATTTG CCGGTGTCGC GCTGCTGATG GCGGCGCGTG
      401     CGGAAGAGGG CCGCGAAGTC GGCTGGTTTC GCTGCCTGCT GGTGTTGTTG
45     451     CCGGCGCGCG GCTTTGTGTC CGCTATGCGT CCGACGCAAA GGCTGATTGC
      501     ACCCATCGGC GCACCGGCAT TCACATCTGT TTCCATTGCC GCCGCATCGT
25     551     TGATGTGCCT GCCGTTTTCG CTTGCTTTGG CGCAAAGTTA TACCGTGGAC
      601     TGGAGCGTCG GAATGGTATT GTCGCTGCTG TATTGGGCG TGGGCTGCAG
      651     CTGGTACGCC TATTGGCTGT GGAACAAGGG GATGAGCCGT GTTCTGCCA
      701     ACGTTTCGGG ACTGTTGATT TCGCTCGAAC CCGTCGTCGG CGTGCTGCTG
      751     GCGGTTTTGA TTTTGGGCGA ACACCTGTCT CCCGTGTCCG TCTTGGGCGT
30     801     GTTTGTCGTC ATCGCCGCCA CCTTGGTTGC CGGCCGGCTG TCGCATCAA
      851     AATAA

```

This encodes a protein having amino acid sequence <SEQ ID 246>:

```

1      MFYQILALII WSSSFIAAKY VYGGIDPALM VGVRLIIAAL PALPACRRHV
35     51      GKIPREEWKP LLIVSFVNYV LTLQLQFVGL KYTSAASASV IVGLEPLLMV
      101     FVGHHFFNDK ARAYHWICGA AAFAGVALLM AGGAEEGGEV GWFGLLVLL
      151     AGAGFCAAMR PTQRLIARIG APAFTSVSIA AASLMCLPFS LALAQSYTVD
      201     WSVGMVLSLL YLGVGCSWYA YWLWNKGMSR VPANVSGLLI SLEPVVGVL
251     AVLILGEHLS PVSVLGVFV IAATLVAGRL SHQK*

```

ORF62a and ORF62-1 show 98.9% identity in 284 aa overlap:

```

40     orf62a.pep MFYQILALIIWSSSFIAAKYVYGGIDPALMVGVRLLIAALPALPACRRHV GKIPREEWKP 60
      orf62-1    MFYQILALIIWSSSFIAAKYVYGGIDPALMVGVRLLIAALPALPACRRHV GKIPREEWKP 60

45     orf62a.pep LLIVSFVNYVLTLLQLQFVGLKYTSAASASVIVGLEPLLMVFVGHFFNDKARAYHWICGA 120
      orf62-1    LLIVSFVNYVLTLLQLQFVGLKYTSAASASVIVGLEPLLMVFVGHFFNDKARAYHWICGA 120

50     orf62a.pep AAFAGVALLMAGGAEEGGEVGVFGCLLVLLAGAGFCAAMRPTQRLIARIGAPAFTSVSIA 180
      orf62-1    AAFAGVALLMAGGAEEGGEVGVFGCLLVLLAGAGFCAAMRPTQRLIARIGAPAFTSVSIA 180

55     orf62a.pep AASLMCLPFSLALAQSYTVDWSVGMVLSLLYLGVGCSWYAYWLWNKGMSRVPANVSGLLI 240
      orf62-1    AASLMCLPFSLALAQSYTVDWSVGMVLSLLYLGLGCGWYAYWLWNKGMSRVPANVSGLLI 240

      orf62a.pep SLEPVVGVLAVLILGEHLSPVSVLGVFVVIATLVAGRLSHQKX 285
      orf62-1    SLEPVVGVLAVLILGEHLSPVSVLGVFVVIATLVAGRLSHQKX 285

```

60 Homology with a predicted ORF from *N.gonorrhoeae*

ORF62 shows 99.5% identity over a 216aa overlap with a predicted ORF (ORF62.ng) from *N. gonorrhoeae*:

	orf62.pep	MFYQILALIIWSSSFIAAKYVYGGIDPALMVGVRLIIAALPALPACRRHVHGKIPREEWKP	60
	orf62ng	MFYQILALIIWSSSFIAAKYVYGGIDPALMVGVRLIIAALPALPACRRHVHGKIPREEWKP	60
5	orf62.pep	LLIVSFVNYVLTLLQLQFVGLKYTSAASASVIVGLEPLLMVFVGHFFFNDKARAYHWICGA	120
	orf62ng	LLIVSFVNYVLTLLQLQFVGLKYTSAASASVIVGLEPLLMVFVGHFFFNDKARAYHWICGA	120
10	orf62.pep	AAFAGVALLMAGGAEEGGEVGVWFGCLLVLLAGAGFCAAMRPTQRLIARIGAPFTSVSIA	180
	orf62ng	AAFAGVALLMAGGAEEGGEVGVWFGCLLVLLAGAGFCAAMRPTQRLIARIGAPFTSVSIA	180
	orf62.pep	AASLMCLPFSLALAQSYTVDWSVGMVLSLLYLGLGC	216
15	orf62ng	AASLMCLPFSLALAQSYTVDWSVGMVLSLLYLGLGCGWYAYWLWNKGMSRVPANASGLLI	240

The complete length ORF62ng nucleotide sequence <SEQ ID 247> is:

	1	ATGTTTACC	AAATCCTGC	CCTGATTATC	TGGGGCAGCT	CGTTTATTGC
	51	CGCCAAATAT	GTCTATGGCG	GCATCGATCC	CGCATTGATG	GTCGGCGTGC
20	101	GCCTGCTGAT	TGCCGCGCTG	CCTGCACTGC	CCGCCGCGCG	CCGTCAATGC
	151	GGCAAGATTG	CGCGTGAGGA	ATGGAAGCCG	TTGCTGATTG	TGTCGTTCTG
	201	CAACTATGTG	CTGACCTCTG	TGCTTCAGTT	TGTCGGGTTG	AAATACACTT
	251	CGCCGCGCAG	CGCATCGGTC	ATTGTGCGAC	TCGAGCCGCT	GCTGATGGTG
	301	TTTGTGCGAC	ACTTTTCTT	CAACGACAAA	GCGCGTGCCT	ACCACTGGAT
25	351	ATGCGGCGCG	GCGGCATTTC	CCGGTGTGCG	GCTGCTGATG	GCGGGCGGTG
	401	CGGAAGAGGG	CGCGGAAGTC	GGCTGGTTCG	GCTGCCTGCT	GGTGTGTGTT
	451	GCGGGCGCGG	GCTTTTGTGC	CGCTATGCGT	CCGACGCAAA	GGCTGATTGC
	501	CCGCATCGGC	GCACCGGCAT	TCACATCTGT	TTCCATTGCC	GCCGCATCGT
	551	TGATGTGCCT	GCCGTTTTCG	CTTGCTTTGG	CGCAAAGTTA	TACCGTGGAC
30	601	TGGAGCGTCG	GGATGGTATT	GTCGCTGTTG	TATTTGGGTT	TGGGGTGCGG
	651	CTGGTACGCC	TATTGGCTGT	GGAACAAGGG	GATGAGCCGT	GTTCCTGCCA
	701	ACGCGTCGGG	ACTGTTGATT	TCGCTCGAAC	CCGTCGTCGG	CGTGCTGTTG
	751	GCGGTTTGA	TTTTGGGCGA	ACATTTATCG	CCCGTGTCGG	CCTTGGGCGT
	801	GTTTGTGCTC	ATCGCCGCCA	CTTTCGCCGC	CGGCCGGCTG	TCGCGCAGGG
	851	ACGCGCAAAA	CGGCAATGCC	GTCTGA		

35 This encodes a protein having amino acid sequence <SEQ ID 248>:

	1	<u>MFYQILALII</u>	<u>WGSSSFIAAKY</u>	<u>VYGGIDPALM</u>	<u>VGVRLLIAAL</u>	<u>PALPACRRHV</u>
	51	<u>GKIPREEWKP</u>	<u>LLIVSFVNYV</u>	<u>LTLLQLQFVGL</u>	<u>KYTSASASV</u>	<u>IVGLEPLLMV</u>
40	101	<u>EVGHFFFNK</u>	<u>ARAYHWICGA</u>	<u>AAFAGVALLM</u>	<u>AGGAEEGGEV</u>	<u>GWFGCLLVLL</u>
	151	<u>AGAGFCAAMR</u>	<u>PTQRLIARIG</u>	<u>APAFTSVSIA</u>	<u>AASLMCLPFS</u>	<u>LALAQSYTVD</u>
	201	<u>WSVGMVLSLL</u>	<u>YLGLGCGWYA</u>	<u>YWLWNKGMSR</u>	<u>VPANASGLLI</u>	<u>SLEPVVGVL</u>
	251	<u>AVLILGEHLS</u>	<u>PVSALGVFVV</u>	<u>IAATFAAGRL</u>	<u>SRRDAQNGNA</u>	<u>V*</u>

ORF62ng and ORF62-1 show 97.9% identity in 283 aa overlap:

		10	20	30	40	50	60
45	orf62ng.pep	MFYQILALIIWSSSFIAAKYVYGGIDPALMVGVRLIIAALPALPACRRHVHGKIPREEWKP					
	orf62-1	MFYQILALIIWSSSFIAAKYVYGGIDPALMVGVRLIIAALPALPACRRHVHGKIPREEWKP					
		10	20	30	40	50	60
50	orf62ng.pep	LLIVSFVNYVLTLLQLQFVGLKYTSAASASVIVGLEPLLMVFVGHFFFNDKARAYHWICGA					
	orf62-1	LLIVSFVNYVLTLLQLQFVGLKYTSAASASVIVGLEPLLMVFVGHFFFNDKARAYHWICGA					
		70	80	90	100	110	120
55	orf62ng.pep	AAFAGVALLMAGGAEEGGEVGVWFGCLLVLLAGAGFCAAMRPTQRLIARIGAPFTSVSIA					
	orf62-1	AAFAGVALLMAGGAEEGGEVGVWFGCLLVLLAGAGFCAAMRPTQRLIARIGAPFTSVSIA					
		130	140	150	160	170	180
60	orf62ng.pep	AASLMCLPFSLALAQSYTVDWSVGMVLSLLYLGLGCGWYAYWLWNKGMSRVPANASGLLI					
	orf62-1	AASLMCLPFSLALAQSYTVDWSVGMVLSLLYLGLGCGWYAYWLWNKGMSRVPANASGLLI					
		190	200	210	220	230	240
65	orf62ng.pep	AASLMCLPFSLALAQSYTVDWSVGMVLSLLYLGLGCGWYAYWLWNKGMSRVPANASGLLI					
	orf62-1	AASLMCLPFSLALAQSYTVDWSVGMVLSLLYLGLGCGWYAYWLWNKGMSRVPANASGLLI					
		190	200	210	220	230	240

-180-

```

                250      260      270      280      290
orf62ng.pep    SLEPVVGVLAVLILGEHLSFVSALGVFVVAATFAAGRLSRRDQNGNAVX
                |||||||:|||||:
5  orf62-1      SLEPVVGVLAVLILGEHLSFVSALGVFVVAATLVAGRLSHQKX
                250      260      270      280

```

Furthermore, ORF62ng shows significant homology to a hypothetical *H.influenzae* protein:

```

sp|Q57147|Y976_HAEIN HYPOTHETICAL PROTEIN HI0976 >gi|1074589|pir||B64163
hypothetical protein HI0976 - Haemophilus influenzae (strain Rd KW20)
10 >gi|1574004 (U32778) hypothetical [Haemophilus influenzae] Length = 128
    Score = 106 bits (262), Expect = 2e-22
    Identities = 56/114 (49%), Positives = 68/114 (59%)

Query: 1  MFYQILALIIWGSSFIAAKYVYGGIDPALMVGVRXXXXXXXXXXCRRHVGVKIPREEWKP 60
15 M YQILAL+IW SS I K Y +DP L+V VR R KI + K
Sbjct: 1  MLYQILALLIWSSSLIVGKLTYSMMDPVLVQVRLIAMIIVMPLFLRRWKIDKPMRKQ 60

Query: 61  LLIVSFVNYVLTLLQFVGLKYTSAASASVIVGLEPLLVMFVGHHFFNDKARAY 114
20 L ++F NY LLQF+GLKYTSA+SA ++GLEPLL+VFGVHFFF K +
Sbjct: 61  LWWLAFFNYTAVFLLQFIGLKYTSASSAVTMIGLEPLLVMFVGHHFFKTKQNGF 114

```

Based on this analysis, including the homology with the transmembrane protein of *H.influenzae* and the putative leader sequence and several transmembrane domains in the gonococcal protein, it is predicted that these proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 30

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 249>:

```

1  ATGCGCCGTT TTCTACCGAT CGCAGCCATA TCGCGmGwms TCCTGkkGTA
51  sGGACTGACG GCGGCAACCG GCAGCACCAG TTCGCTGGCG GATTATTTCT
30 101 GGTGGATTGT TCGTTCAGC GCAATGCTGC TGCTGGTGT GTCCGCCGTT
151 TTGGCACGTT ATGTCATATT GCTGTTGAAA GACAGGCGCG ACGGCGTATT
201 CCGTTCGCTA srTyGCCAAA gsGCTGkks TGGG.ATGTT TACGCTGGTT
251 GCCGkACTGC CCGGCGTGTT TCTGTTGCGC TTTCCCGCAC AGTTCATCAA
301 CGGCACGATT AATTCGTGGT TCGGCAACGA TACCCACGAG GCGCTTGAAC
35 351 GCAGCCTCAA TTTGAGCAAG TCCGCATTGA ATTTGGCGGC AGACAACGCC
401 CTCGGCAACG CCGTCCCCGT GCAGATAGAC CTCATCGGCG CGGCTTCCCT
451 GCCCGGGGAT ATGGGCAGGG TGCTGGAACA TTACGCCGCG AGCGGTTTTG
501 CCCAGCTTGC CCTGTACAAY ksCGCAACG GCAAAATCGA AAAAAGCATC
551 AACCAGCACA AGCTCGATCA GCCGTTTCCA GGTAAGGCGC GTTGGGAaAa
40 601 AATCCaACGG GCGGGTTCGG TCAGGGATTG GGAAGCATA GCGGCGGTAT
651 TGTaCGCGCA GGGCTGGCTG TCGGCGGGTA CGCACwACGG GCGCGATTAC
701 GCCTTGTTTT TCCGTCAGCC GGTTCCTAAA GCGGTGGCAG AGGATGCCGT
751 YTTAATCGAA AAGGCAAGGG CGAAATATGC TGAGTTGAGT TACAGCAAAA
801 AAGGTTTGCA GACCTTTTTT CTGGCAACCC TGCTGATTGC CTCGCTGCTG
45 851 TCGATTTTTT TTGCACTGGT CATGGCACTG TATTTCGCCC GCGGTTTCGT
901 CGAACCCGTC CTATCGCTTG CCGAGGGGGC GAAGGCGGTG GCGCAAGGGC
951 ATTTACAGCA GACGCGCCCC GTGTTGCGCA ACGACGAGT CGGACGCTTG
1001 ACCArGTTGT TCAACCACAT GACCGAGCAG CTTTCCATCG CCAAAGATGC
1051 AGACGAGCGC AACCGCCGGC GCGAGGAAGC CGCCAGGCAT TATCTTGAAT
50 1101 GCGTGTGGA GGGGCTGACC ACGGGCGTGG TGGTGTGTTGA CGAACAAGGC
1151 TGTCTGAAAA CCTTCAACAA AGCGGCGGGT ACC..

```

This corresponds to the amino acid sequence <SEQ ID 250; ORF64>:

```

1  MRRFLPIAAI CAXLXLXGLT AATGSTSSLA DYFWWIVAFS AMLLLVLSAV
51  LARYVILLK DRRDGVFGSX XAKXPXXMF TLVAXLPGVF LFGFPAQFIN
55 101 GTINSWFGND THEALERSLN LSKSALNLAA DNALGNAPV QIDLIGAASL
151 PQDMGRVLEH YAGSGFAQLA LYNXASGKIE KSINPHKLDQ PFPKGARWEK
201 IQRAGSVRDL ESIGGVLYAQ GWLSAGTHXG RDYALFFRQP VPKGVAEDAV
251 LIEKARAKYA ELSYSKGLQ TFFLATLLIA SLLSIFLALV MALYFARRFV

```

301 EPVLSLAEGA KAVAQGDfsQ TRPVLrNDEF GRLTXLFNHM TEQLSIAKDA
 351 DERNRRREEA ARHYLECVLE GLTTGVVVFD EQGCLKTFNK AAGT..

Further work revealed the complete nucleotide sequence <SEQ ID 251>:

```

      1 ATGCGCCGTT TTCTACCGAT CGCAGCCATA TCGCGCGTCG TCCTGTTGTA
5      51 CGGACTGACG GCGGCAACCG GCAGCACCAG TTCGCTGGCG GATTATTTCT
      101 GGTGGATTGT TGCCTTCAGC GCAATGCTGC TGCTGGTGTG GTCCGCCGTT
      151 TTGGCACGTT ATGTCATATT GCTGTGAAA GACAGGCGCG ACGGCGTATT
      201 CGGTTCGCAG ATTGCCAAAC GCCTTTCTGG GATGTTTACG CTGGTTGCCG
      251 TACTGCCCGG CGTGTTCCTG TTCGCGGTTT CCGCACAGTT CATCAACGGC
10     301 ACGATTAATT CGTGGTTCGG CAACGATACC CACGAGGCGC TTGAACGCAG
      351 CCTCAATTG AGCAAGTCCG CATTGAATTT GGCGGCAGAC AACGCCCTCG
      401 GCAACGCCGT CCCCCTGCAG ATAGACCTCA TCGGCGCGGC TTCCCTGCCC
      451 GGGGATATGG GCAGGGTGCT GGAACATTAC GCCGCGAGCG GTTTTGCCCA
      501 GCTTGCCCTG TACAATGCCG CAAGCGGCAA AATCGAAAAA AGCATCAACC
15     551 CTGCACAAGCT CGATCAGCCG TTTCAGGTA AGGCGCGTTG GGAAAAAATC
      601 CAACGGGCGG GTTCGGTCAG GGATTGGAA AGCATAGGCG GCGTATTGTA
      651 CGCGCAGGGC TGGCTGTCGG CGGGTACGCA CAACGGGCGC GATTACGCCT
      701 TGTTTTTCCG TCAGCCGCTT CCCAAGGCG TGGCAGAGGA TGCCGCTCTA
      751 ATCGAAAAGG CAAGGGCGAA ATATGCTGAG TTGAGTTACA GAAAAAAGC
20     801 TTTGCAGACC TTTTCTCTGG CAACCTGCTG GATTGCCTCG CTGCTGTCGA
      851 TTTTCTTTCG ACTGGTCATG GCACGTGATT TCGCCCGCGC TTTCTGCGAA
      901 CCCGTCTTAT CGCTTGCCGA GGGGGCGAAG GCGGTGGCGC AAGGCGATT
      951 CAGCCAGACG CGCCCCGTGT TCGCAACGA CGAGTTCGGA CGCTTGACCA
25    1001 AGTTGTTCAA CCACATGACC GAGCAGCTTT CCATCGCCAA AGAAGCAGAC
      1051 GAGCGCAACC GCCGCGCGCA GGAAGCCGCC AGGCATTATC TTGAATGCGT
      1101 GTTGGAGGGG CTGACCACGG GCGTGGTGGT GTTTGACGAA CAAGGCTGTC
      1151 TGAAAACCTT CAACAAAGCG GCGGAACAGA TTTTGGGGAT GCCGCTTACC
      1201 CCCCTGTGGG GCAGCAGCCG GCACGGTTGG CACGGCGTTT CGGCGCAGCA
      1251 GTCCCTGCTT GCCGAAGTGT TTGCCGCCAT CGGCGCGGCG GCAGGTACGG
30    1301 ACAAACCGCT CCATGTGAAA TATGCCGCGC CGGACGATGC CAAAATCCTG
      1351 CTGGGCAAGG CAACCGTCCT GCCGAAGAC AACGGCAACG GCGTGGTAAT
      1401 GGTGATTGAC GACATCACCG TTTTGATACA CGCGCAAAA GAAGCCGCGT
      1451 GGGGCGAAGT GCGGAAGCGG CTGGCACACG AAATCCGCAA TCCGCTCAGC
      1501 CCCATCCAGC TTTCCGCGCA ACGGCTGGCG TGGAAATTGG GCGGGAAGCT
35    1551 GGATGAGCAG GATGCGCAAA TCCTGACGCG TTCGACCGAC ACCATCGTCA
      1601 AACAGTGGC GGCATTGAAG GAAATGGTCG AAGCATTCGG CAATTATGCG
      1651 CGTTCCCTCT CGCTCAAATT GGAAATCAG GATTTGAACG CCTTAATCGG
      1701 CGATGTGTTG GCATTGTATG AAGCCGGTCC GTGCCGTTT GCGGCGGAGC
      1751 TTGCCGCGCA ACCGCTGACG GTGGCGGCGG ATACGACCGC CATGCCGCG
40    1801 GTGCTGCACA ATATTTTCAA AAATGCCGCC GAAGCGGCGG AAGAAGCCGA
      1851 TGTGCCCGAA GTCAGGGTAA AATCGGAAAC AGGGCAGGAC GGTCCGATTG
      1901 TCTGACGGT TTGCGACAAC GGCAAAGGTT TCGGCAGGGA AATGCTGCAC
      1951 AACGCCTTCG AGCCGTATGT AACGGACAAA CCGGCGGGAA CGGGATTGGG
      2001 TCTGCCTGTG GTGAAAAAAA TCATTGAAGA ACACGCGCGC CGCATCAGCC
45    2051 TGAGCAATCA GGATCGCGGT GCGCGGTGTG TCAGAATCAT CTTGCCAAAA
      2101 ACGGTAAAAA CTTATGCGTA G

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This corresponds to the amino acid sequence <SEQ ID 252; ORF64-1>:

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      1 MRRFLPIAAI CAVVLLYGLT AATGSTSSLA DYFWWIVAFS AMLLLVLSAV
50     51 LARYVILLK DRRDGVFGSQ IAKRLSGMFT LVAVLPGVFL FGVSAQFING
      101 TINSWFGNDT HEALERSLNL SKSALNLAAD NALGNVVPVQ IDLIGAASLP
      151 GDMGRVLEHY AGSGFAQLAL YNAASGKIEK SINPHKLDQP FPGKARWEKI
      201 QRAGSVRDLE SIGGVLYAQG WLSAGTHNGR DYALFFRQPV PKGVAEDAVL
      251 IEKARAKYAE LSYSKKGLQT FFLATLLIAS LLSIFLALVM ALYFARRFVE
55    301 PVLSLAEGAK AVAQGDfsQT RPVLrNDEFG RLTKLFNHMT EQLSIAKEAD
      351 ERNRRREEAA RHYLECVLEG LTTGVVVfDE QGCLKTFNKA AEQILGMPLT
      401 PLWGSSRHGW HGVSAQqSLL AEVFAAIGAA AGTDKPVHVK YAAPDDAKIL
      451 LGKATVLPED NGNGVVMVID DITVLIHAQK EAAWGEVAKR LAHEIRNPLT
      501 PIQLSAERLA WKLGGKLDEQ DAQILTRSTD TIVKQVAALK EMVEAFRNVA
      551 RSPSLKLENQ DLNALIGDVL ALYEAGPCRf AAELAGEPLT VAADTTAMRQ
60    601 VLHNIIFKNA EAAEEADVPE VRVKSETGQD GRIVLTVCND KGKFGREMLH
      651 NAFEPYVTDK PAGTGLGLPV VKKIIIEHGG RISLSNQDAG GACVRIILPK
      701 TVKTYA*

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Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF64 shows 92.6% identity over a 392aa overlap with an ORF (ORF64a) from strain A of *N.*

meningitidis:

5	orf64.pep	10 20 30 40 50 60	MRRFLPIAAICAXLXXGLTAATGSTSSLADYFWWIVAFSAMLVLSAVLARYVILLK
	orf64a	10 20 30 40 50 60	MRRFLPIAAICAVVLLYGLTAATGSTSSLADYFWWIVAFSAMLVLSAVLARYVILLK
10	orf64.pep	70 80 90 100 110 120	DRRDGVFGSXXAKXFXMFTLVAXLPGVFLFGFPAQFINGTINSWFGNDTHEALERSLN
	orf64a	70 80 90 100 110 120	DRRDGVFGSQIAKR-LSGMFTLVAVLPGVFLFGVSAQFINGTINSWFGNDTHEALERSLN
15	orf64.pep	130 140 150 160 170 180	LSKSALNLAADNALGNAVVPQIDLIGAASLPGDMGRVLEHYAGSGFAQLALYNXASGKIE
	orf64a	120 130 140 150 160 170	LSKSALNLAADNALGNAIPQIDIXIGAASLPXDMGRVLEHYAGSGFAQLALYNAASGKIE
20	orf64.pep	190 200 210 220 230 240	KSINPHKLDQFPFGKARWEKIQRAGSVRDLESIGGVLYAQGWLASAGTHXGRDYALFFRQP
	orf64a	180 190 200 210 220 230	KSINPHKLDQFPFGKARWEKIQAGSVRDLESIGGVLYAXGWLASAGTHNGRDYALFFRQP
30	orf64.pep	250 260 270 280 290 300	VPKGVAEDAVLIEKARAKYAELSYSKKGLQTFFLATLLIASLLSIFLALVMALYFARRFV
	orf64a	240 250 260 270 280 290	VPKGVAEDAVLIEKARAXXXLSYSKKGLQTFFLATLLIASLLSIFLALVMALYFARRFV
35	orf64.pep	310 320 330 340 350 360	EPVLSLAEGAKAVAQGDFSQTRPVLNRNDEFGRLTXLFNHMTQLSIAKDADERNRRREEA
	orf64a	300 310 320 330 340 350	EPVLSLAEGAKAVAQGDFSQTRPVLNRNDEFGRLTKLFNHMTQLSIAKEADERNRRREEA
40	orf64.pep	370 380 390	ARHYLECVLEGLTTGVVVFDEQGCLKTFNKAAGT
	orf64a	360 370 380 390 400 410	ARHYLECVLEGLTTGVVVFDEQGCLKTFNKAAEQILGMPLTPLWGSSSRHGWGVSAQQSL
45	orf64a	420 430 440 450 460 470	LAEVFAAIGAAAGTDKPVHVKYAAPDDAKILLGKATVLPEDNXNGVVMIDDITVLIHAQ

The complete length ORF64a nucleotide sequence <SEQ ID 253> is:

50	1	ATGCGCCGTT	TTCTACCGAT	CGCAGCCATA	TGCGCCGTCG	TCCTGTTGTA
	51	CGGACTGACG	GCGGCAACCG	GCAGCACCAG	TTCGCTGGCG	GATTATTTC
	101	GGTGGATTGT	TGCGTTCAGC	GCAATGCTGC	TGCTGGTGTT	GTCCGCCGTT
	151	TTGGCACGTT	ATGTCATATT	GCTGTTGAAA	GACAGGCGCG	ACGGCGTATT
	201	CGGTTTCGAG	ATTGCCAAAC	GCCTTTCCGG	GATGTTTACG	CTGGTTGCCG
55	251	TACTGCCCGG	CGTGTTCCTG	TTCGGCGTTT	CCGCACAGTT	TATCAACGGC
	301	ACGATTAATT	CGTGGTTCGG	CAACGATACC	CACGAGGCGC	TGAACGCAG
	351	CCTCAATTG	AGCAAGTCCG	CATTGAATCT	GGCGGCAGAC	AACGCCCTTG
	401	GCAACGCCAT	CCCCGTGCAG	ATAGACNTCA	TCGGCGGCGC	TCCTTGCCC
	451	NGGGATATGG	GCAGGGTGCT	GGAACATTAC	GCCGGCAGCG	GTTTGTGCCA
60	501	GCTTGCCCTG	TACAATGCCG	CAAGCGGCAA	AATCGAAAAA	AGCATCAACC
	551	CGCACAAAGCT	CGATCAGCCG	TTTCCAGGTA	AGGCGCGGTT	GGAAAAAATC
	601	CAACAGGCGG	GTTCGGTCAG	GGATNNGGAA	AGCATAGGCG	GCGTATTGTA
	651	CGCGCANGGC	TGGCTGTCGG	CAGNNACGCA	CAACGGGCGC	GATTACGCCT
	701	TGTTTTTCCG	TCAGCCGCTT	CCCAAAGGCG	TGGCAGAGGA	TGCCGTCTTA
65	751	ATCGAAAAGG	CAAGGGCGNA	ANANNNTNAG	TTGAGTTACA	GCAAAAAAGG
	801	TTTGCAGACC	TTTTCCTTNG	CAACCCTGCT	GATTGCCTCN	CTGCTGTCGA
	851	TTTTTCTTGC	ACTGGTCATG	GCACTGTATT	TCGCCCGCCG	TTTCGTCGAA

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901 CCCGTCCTAT CGCTTGCCGA GGGGGCGAAG GCGGTGGCGC AAGGCGATTT
951 CAGCCAGACG CGCCCCGTGT TGCGCAACGA CGAGTTCGGA CGCTTGACCA
1001 AGTTGTTCAA CCACATGACC GAGCAGCTTT CCATCGCCAA AGAAGCAGAC
1051 GAGCGCAACC GCCGGCGCGA GGAAGCCGCC AGACATTATC TCGAATGCGT
1101 GTTGGAGGGG CTGACCACGG GCGTGGTGGT GTTTGACGAA CAAGGCTGTC
1151 TGAAAACCTT CAACAAAGCG GCGGAACAGA TTTTGGGGAT GCCGCTTACC
1201 CCCCTGTGGG GCAGCAGCCG GCACGGTTGG CACGGCGTTT CGGCGCAGCA
1251 GTCCCTGCTT GCCGAAGTGT TTGCCGCCAT CGGCGCGGCG GCAGGTACGG
1301 ACAAACCGGT CCATGTGAAA TATGCCGCGC CGGACGATGC CAAAATCCTG
1351 CTGGGCAAGG CAACCGTCCT GCCCGAAGAC AACNGCAACG GCGTGGTAAT
1401 GGTGATTGAC GACATCACCG TTTTGATACA CGCGCAAAA GAAGCCGCGT
1451 GGGGCGAAGT GGCAAAACGG CTGGCACACG AAATCCGCAA TCCGCTCAGC
1501 CCCATCCAGC TTTCTGCCGA ACGGCTGGCG TGGAAATTGG GCGGGAAGCT
1551 GGACGAGCAN GACGCGCAAA TCCTGACACG TTCGACCGAC ACCATCATCA
1601 AACAAAGTGG GCATTAAAA GAAATGGTCG AGGCATTCCG CAATTACNCG
1651 CGTTCCCTTT CGNCTCAATT GGAAAATCAG GATTTGAACG CCTTAATCGG
1701 CGATGTGTTG AAGCTGTACG AAGCTGGTCC GTGCCGTTT GCGGCGGAAC
1751 TTGCCGGCGA ACCGCTGATG ATGGCGGCGG ATACGACCGC CATGCGGCAG
1801 GTGCTGCACA ATATTTTCAA AAATGCCGCC GAAGCGGCGG AAGAAGCCGA
1851 TGTGCCCGAA GTCAGGGTAA AATCGGAAGC GGGGCAGGAC GGACGGATTG
1901 TCCTGACAGT TTGCGACAAC GGCAAGGGGT TCGGCAGGGA AATGCTGCAC
1951 AATGCCTTCG AGCCGTATGT AACGGACAAA CCGGCTGGAA CGGGATTGNG
2001 ACTGCCCCGT GTGAAAAAAA TCATTGAAGA ACACGGCGGC CNCATCAGCC
2051 TGAGCAATCA GGATGCGGGC GGCGCGTNTG TCAGAATCAT CTTGCCAAAA
2101 ACGGTAGAAA CTTATGCGTA G

```

This encodes a protein having amino acid sequence <SEQ ID 254>:

30
35
40

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1 MRRFLPIAAI CAVVLLYGLT AATGSTSSLA DYFWWIVAFS AMLLLVLSAV
51 LARYVILLLK DRRDGVFGSQ IAKRLSGMFT LVAVLPGVFL FGVSAQFING
101 TINSWFGNDT HEALERSLNL SKSALNLAAD NALGNAIPVQ IDXIGAASLP
151 XDMGRVLEHY AGSGFAQLAL YNAASGKIEK SINPHKLDQP FPGKARWEKI
201 QQAGSVRDXE SIGGVLYXG WLSAXTHNGR DYALFFRQPV PKGVAEDAVL
251 IEKARAXXXX LYSKKGLQT FFLATLLIAS LLSIFLALVM ALYFARRFVE
301 PVLSLAEGAK AVAQGDFSQT RPLVRNDEFG RLTKLFNHMT EQLSIAKEAD
351 ERNRRREEAA RHYLECVLEG LTTGVVVFDE QGCLKTFNKA AEQILGMPLT
401 PLWGSSRHGW HGVSAAQSL L AEFVFAAIGAA AGTDKPVHVK YAAPDDAKIL
451 LGKATVLPED NXNGVVMVID DITVLIHAQK EAAWGEVAKR LAHEIRNPLT
501 PIQLSAERLA WKLGGKLDX DAQILTRSTD TTIKQVAALK EMVEAFRNYX
551 RSPSXQLENQ DLNALIGDVL ALYEAGPCRF AAELAGEPLM MAADTTAMRQ
601 VLHNIKNAE EAAEEADVPE VRVKSEAGQD GRIVLTVCDN KGKFGREMLH
651 NAFEPYVTDK PAGTGLXLPV VKKIEEHGG XISLSNQDAG GAXVRIILPK
701 TVETYA*

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ORF64a and ORF64-1 show 96.6% identity in 706 aa overlap:

45
50
55
60
65

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              10      20      30      40      50      60
orf64a.pep  MRRFLPIAAICAVVLLYGLTAATGSTSSLADYFWWIVAFSAML LLLVLSAVLARYVILLK
|||||
orf64-1      MRRFLPIAAICAVVLLYGLTAATGSTSSLADYFWWIVAFSAML LLLVLSAVLARYVILLK
              10      20      30      40      50      60

              70      80      90      100     110     120
orf64a.pep  DRRDGVFGSQIAKRLSGMFTLVAVLPGVFLFGVSAQFINGTINSWFGNDTHEALERSLNL
|||||
orf64-1      DRRDGVFGSQIAKRLSGMFTLVAVLPGVFLFGVSAQFINGTINSWFGNDTHEALERSLNL
              70      80      90      100     110     120

              130     140     150     160     170     180
orf64a.pep  SKSALNLAADNALGNAIPVQIDXIGAASLPXDMGRVLEHYAGSGFAQLALYNAASGKIEK
|||||
orf64-1      SKSALNLAADNALGNAIPVQIDLIGAASLPXDMGRVLEHYAGSGFAQLALYNAASGKIEK
              130     140     150     160     170     180

              190     200     210     220     230     240
orf64a.pep  SINPHKLDQFPFGKARWEKIQAGSVRDLESIGGVLYXGWLSAXTHNGRDYALFFRQPV
|||||
orf64-1      SINPHKLDQFPFGKARWEKIQAGSVRDLESIGGVLYXGWLSAXTHNGRDYALFFRQPV
              190     200     210     220     230     240

              250     260     270     280     290     300

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5	orf64a.pep	PKGVAEDAVLIEKARAXXXLSYSKKGLQTFFLATLLIASLLSIFLALVMALYFARRFVE
	orf64-1	PKGVAEDAVLIEKARAKYAELSYSKKGLQTFFLATLLIASLLSIFLALVMALYFARRFVE
10	orf64a.pep	FVLSLAEGAKAVAQGDFSQTRFVLRNDEFGRITKLFNHEQLSIAKEADERNRRREEAA
	orf64-1	FVLSLAEGAKAVAQGDFSQTRFVLRNDEFGRITKLFNHEQLSIAKEADERNRRREEAA
15	orf64a.pep	RHYLECVLEGLTTGVVVFDEQGCLKTFNKAAEQILGMPLTPLWGSSRHGWHGVSAQQSLL
	orf64-1	RHYLECVLEGLTTGVVVFDEQGCLKTFNKAAEQILGMPLTPLWGSSRHGWHGVSAQQSLL
20	orf64a.pep	AEVFAAIGAAAGTDKPVHVKYAAPDDAKILLGKATVLPEDNXNGVVMVDDITVLIHAQK
	orf64-1	AEVFAAIGAAAGTDKPVHVKYAAPDDAKILLGKATVLPEDNGNGVVMVDDITVLIHAQK
25	orf64a.pep	EAAWGEVAKRLAHEIRNPLTPIQLSAERLAWKLGKGLDEXDAQILTRSTDITIKQVAALK
	orf64-1	EAAWGEVAKRLAHEIRNPLTPIQLSAERLAWKLGKGLDEQDAQILTRSTDITIVKQVAALK
30	orf64a.pep	EMVEAFRNYXRSXQLENQDNLALIGDVLALYEAGPCRFAAELAGEPLMMAADTTAMRQ
	orf64-1	EMVEAFRNYARSPSLKLENQDNLALIGDVLALYEAGPCRFAAELAGEPLTVAADTTAMRQ
35	orf64a.pep	VLHNI FKNAAEAAEEADVPEVRVKSEAGQDGRIVLTVCNKGKFGREMLHNAFEPYVTDK
	orf64-1	VLHNI FKNAAEAAEEADVPEVRVKSETGQDGRIVLTVCNKGKFGREMLHNAFEPYVTDK
40	orf64a.pep	PAGTGLXLPVVKKIIIEHGGXISLSNQDAGGAXVRIILPKTVETYAX
	orf64-1	PAGTGLGLPVVKKIIIEHGGRIISLSNQDAGGACVRIILPKTVKTYAX

Homology with a predicted ORF from *N.gonorrhoeae*ORF64 shows 86.6% identity over a 387aa overlap with a predicted ORF (ORF64.ng) from *N.*50 *gonorrhoeae*:

55	orf64.pep	MRRFLPIAAICAXLXXGLTAATGSTSSLADYFWWIVAFSAML LLVL SAVLARYVILLK	60
	orf64ng	MRRFLPIAAICAVVLLYGLTAATGSTSSLADYFWWIVSFSAML LLVL SAVLARYVILLK	60
60	orf64.pep	DRRDGVFGSXXAKXPXXMFTLVAXLPGVFLFGFPAQFINGTINSWFGNDTHEALERSLN	120
	orf64ng	DRRNGVFGSQIAKR-LSGMFTLVAVLPGLFLFGISAQFINGTINSWFGNDTHEALERSLN	119
65	orf64.pep	LSKSALNLAADNALGNVAVPVQIDLIGAASLPGDMGRVLEHYAGSGFAQLALYNXASGKIE	180
	orf64ng	LSKSALDLAADNAVSNVAVPVQIDLIGTASLSGNMGSVLEHYAGSGFAQLALYNAASGKIE	179
65	orf64.pep	KSINPHKLDQPFPGKARWEKIQRAGSVRDLESIGGVLYAQGWLSAGTHXGRDYALFFRQP	240
	orf64ng	KSINPHQFDQPLPDKEHWEQIQQTGSVRSLESIGGVLYAQGWLSAGTHNGRDYALFFRQP	239

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orf64.pep	VPKGVAEDAVLIEKARAKYAELSYSKKGLQTFFLATLLIASLLSIFLALVMALYFARRFV	300
orf64ng	IPENVAQDAVLIEKARAKYAELSYSKKGLQTFFLVTLIIASLLSIFLALVMALYFARRFV	299
5 orf64.pep	EPVLSLAEGAKAVAQGDFSQTRPVLNRNDEFGRLTXLFNHMTQELSIAKDADERNRRREEA	360
orf64ng	EPILSLAEGAKAVAQGDFSQTRPVLNRNDEFGRLTXLFNHMTQELSIAKEADERNRRREEA	359
10 orf64.pep	ARHYLECVLEGLTTGVVVVFDEQGCCLKTFNKAAGT	394
orf64ng	ARHYLECVLDGLTTGVVVSYPLSCCRTAVFSTCHSSPLSYF	400

An ORF64ng nucleotide sequence <SEQ ID 255> was predicted to encode a protein having amino acid sequence <SEQ ID 256>:

1	MRRFLPIAAI	CAVVLLYGLT	AATGSTSSLA	DYFWWIVSFS	AMLLLVLSAV
15 51	LARYVILLK	DRRNGVFGSQ	IAKRLSGMFT	LVAVLPGLFL	FGISAQFING
101	TINSWFGNDT	HEALERSLNL	SKSALDLAAD	NAVSNAPVQ	IDLIGTASLS
151	GNMGSVLEHY	AGSGFAQLAL	YNAASKIEK	SINPHQFDQP	LPDKEHWEQI
201	QQTGVSRSLE	SIGGVLYAQG	WLSAGTHNGR	DYALFFRQPI	PENVAQDAVL
251	IEKARAKYAE	LSYSKGLQT	FFLVTLIIAS	LLSIFLALVM	ALYFARRFVE
20 301	PILSLAEGAK	AVAQGDFSQT	RPVLRNDEFG	RLTKLFNHMT	EQLSIAKEAD
351	ERNRRREEAA	RHYLECVLDG	LTTGVVVSYP	LSCCRTAVFS	TCHSSPLSYF*

Further work revealed the complete gonococcal DNA sequence <SEQ ID 257>:

1	ATGCGCCGCT	TCCTACCGAT	CGCAGCCATA	TGCGCCGTCG	TCCTGCTGTA
25 51	CGGATTGACG	GCGGCGACCG	GCAGCACCAG	TTCGCTGGCG	GATTATTTCT
101	GGTGGATAGT	CTCGTTCAGC	GCAATGCTGC	TGCTGGTGTT	GTCCGCCGTT
151	TTGGCACGTT	ATGTCATATT	GCTGTTGAAA	GACAGGCGCA	ACGGCGTGTT
201	CGGTTTCGAG	ATTGCCAAAC	GCCTTTCCGG	GATGTTACAG	CTGGTCGCCG
251	TACTGCCCGG	CTTGTTCCCTG	TTCGGCATT	CCGCGCAGTT	TATCAACGGC
301	ACGATTAATT	CGTGGTTCGG	CAACGACACC	CACGAAGCCC	TGCAACGCAG
351	CCTTAATTTG	AGCAAGTCCG	CACTGGATTT	GGCGGCAGAC	AATGCCGTCA
401	GCAACGCCGT	TCCCGTACAG	ATAGACCTCA	TGCGGCACCG	CTCCCTGTCTG
451	GGCAATATGG	GCACTGTGCT	GGAACACTAC	GCCGGCAGCG	GTTTGTGCCA
501	GCTTGCCCTG	TACAATGCCG	CAAGCGGGAA	AATCGAAAAA	AGCATCAATC
551	CGCACCAATT	CGACCAGCCG	CTTCCCGACA	AAGAACATTG	GGAACAGATT
601	CAGCAGACCG	GTTCCGTTTCG	GAGTTTGGA	AGCATAGGCG	GCGTATTGTA
651	CGCGCAGGGA	TGGTTGTCGG	CAGGTACGCA	CAACGGGCGC	GATTACGCGC
701	TGTTCTTCCG	CCAGCCGATT	CCCGAAAATG	TGGCACAGGA	TGCCGTTCTG
751	ATTGAAAAGG	CGCGGGCGAA	ATATGCCGAA	TTGAGTTACA	GCAAAAAGG
801	TTTGACAGAC	TTTTTCTCG	TAACCTTGCT	GATTGCCTCG	CTGCTGTCCA
851	TTTTTCTTGC	GCTGGTAATG	GCACTGTATT	TGCCCCGCG	TTTCGTGCAA
901	CCCATTCTGT	CGCTTGCCGA	GGGCGCAAAG	GCGGTGGCGC	AGGGTGATTT
951	CAGCCAGACG	CGCCCCGTAT	TGCGCAACGA	CGAGTTCCGA	CGTTTGACCA
1001	AGCTGTTCAA	CCATATGACC	GAGCAGCTTT	CCATCGCCAA	AGAAGCAGAC
1051	GAACGCAACC	GCCGGCGCGA	GGAAGCCGCC	CGTCACTACC	TGAGTGCGT
1101	GTTGGATGGG	TTGACTACCG	GTGTGGTGGT	GTTTGACGAA	AAAGGCCGTT
1151	TGAAAACCTT	CAACAAGGCG	GCGGAACAGA	TTTTGGGGAT	GCCGCTCGCC
1201	CCCCTGTGGG	GCAGCAGCCG	GCACGGTTGG	CACGGCGTTT	CGGCGCAGCA
1251	GTCCCTGCTT	GCCGAAGTGT	TtgcgcccAT	CGGTGCGGCG	GCAGGTACGG
1301	ACAAACCGGT	CCAGGTGGAA	TATGCCGCGC	CGGACGATGC	CAAAATCCTG
1351	CTGGGCAAGG	CGACGGTATT	GCCCGAAGAC	AACGGCAACG	GCGTGGTGAT
1401	GGTGATTGAC	GACATCACCG	TGCTGATACG	CGCGCAAAAA	GAAGCCGCGT
1451	GGGGTGAAGT	GGCGAAGCGG	CTGGCACACG	AAATCCGCAA	TCCGCTCACG
1501	CCCATCCAGC	TTTCCGCCGA	ACGGCTGGCG	TGGAATTTGG	GCGGGAAGCT
1551	GGACGATCAG	GACGCGCAAA	TCCTGACGCG	TtcgACCGAC	ACCATCATCA
1601	AACAGgtggc	gGCGTTAAAA	GAAATGGTCG	AGGCATTCCG	CAATTACGCG
1651	CGCGCCCCTT	CGCTCAAAC	GGAAAATCAG	GATTTGAAACG	CCTTAATCGG
1701	CGATGTTTTG	GCCCTGTACG	AAGCCGGCCC	GTGCCGTTT	GAGGCGGAAC
1751	TTGCCGCGGA	ACCGCTGATG	ATGGCGGCGG	ATACGACCGC	CATGCGGCAG
1801	GTGCTGCACA	ATATTTTCAA	AAATGCCGCC	GAAGCGGCGG	AAGAAGCCGA
1851	TATGCCCGAA	GTCAGGTAA	AATCGGAAAC	GGGGCAGGAC	GGACGGATTG
1901	TCCTGACGGT	TTGCGACAAC	GGCAAGGGAT	TCGGCAAGGA	AATGCTGCAC
1951	AATGCTTTTCG	AGCCGTATGT	GACGGATAAG	CCGGCGGGAA	CGGGACTGGG
2001	TCTGCCTGTA	GTGAAAAAAA	TCATTGGAGA	ACACGGCGGC	CGCATCAGCC
2051	TGAGCAATCA	GGATGCGGGT	GGGGCGTGTG	TCAGAATCAT	CTTGCCAAAA
2101	ACGGTAGAAA	CTTATGCGTA	G		

This corresponds to the amino acid sequence <SEQ ID 258; ORF64ng-1>:

```

      1 MRRFLPIAAI CAVVLLYGLT AATGSTSSLA DYFWWIVSFS AMLLLVLSAV
    51 LARYVILLK  DRRNGVFGSQ IAKRLSGMFT LVAVLPGLFL FGISAQFING
   101 TINSWFGNDT HEALERSLNL SKSALDLAAD NAVSNAVVPQ IDLIGTASLS
    151 QMGSVLEHY AGSGFAQLAL YNAASGKIEK SINPHQFDQP LPDKEHWEQI
    201 QQTGSVRSLE SIGGVLYAQQ WLSAGTHNGR DYALFFRQPI PENVAQDAVL
    251 IEKARAKYAE LSYSKKGLQT FFLVTLIIAS LLSIFLALVM ALYFARRFVE
    301 PILSLAEGAK AVAQGDFSQT RPYLRNDEFG RLTKLFNHMT EQLSIAKEAD
    351 ERNRRREEAA RHYLECVLDG LTTGVVVFDE KGRKTFNKA AEQILGMPLA
    401 PLWGSSRHGW HGVSAQQSLL AEVFAAIGAA AGTDKPVQVE YAAPDDAKIL
    451 LGKATVLPED NGNGVVMVID DITVLIRAQK EAAWGEVAKR LAHEIRNPLT
    501 PIQLSAERLA WKLGGKLLDD DAQILTRSTD TTIKQVAALK EMVEAFRNYA
    551 RAPSLKENQ  DLNALIGDVL ALYEAGPCR FEAELAGEPLM MAADTTAMRQ
    601 VLHNIFKNAA EAAEEADMPE VRVKSETGQD GRIVLTVCDN GKGFGKEMLH
    651 NAFEPVYTDK PAGTGLGLPV VKKIIGEHGG RISLSNQDAG GACVRIILPK
    701 TVETYA*
  
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ORF64ng-1 and ORF64-1 show 93.8% identity in 706 aa overlap:

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      10      20      30      40      50      60
    20 orf64ng-1.pep MRRFLPIAAICAVVLLYGLTAATGSTSSLADYFWWIVSFSAMLLLVLSAVLARYVILLK
       orf64-1      MRRFLPIAAICAVVLLYGLTAATGSTSSLADYFWWIVAFSAMLLLVLSAVLARYVILLK
      10      20      30      40      50      60

      70      80      90     100     110     120
    25 orf64ng-1.pep DRRNGVFGSQIAKRLSGMFTLVAVLPGLFLFGISAQFINGTINSWFGNDTHEALERSLNL
       orf64-1      DRRDGVFGSQIAKRLSGMFTLVAVLPGLFVGVSQAQFINGTINSWFGNDTHEALERSLNL
      70      80      90     100     110     120

      130     140     150     160     170     180
    30 orf64ng-1.pep SKSALDLAADNAVSNVQIDLIGTASLSGNMGSVLEHYAGSGFAQLALYNAASGKIEK
       orf64-1      SKSALNLAADNALGNVQIDLIGAASLPGDMGRVLEHYAGSGFAQLALYNAASGKIEK
      130     140     150     160     170     180

      190     200     210     220     230     240
    35 orf64ng-1.pep SINPHQFDQPLPDKEHWEQIQQTGSVRSLESIGGVLYAQQWLSAGTHNGRDYALFFRQPI
       orf64-1      SINPHKLDQPFPGKARWEKIQRAGSVRDLESIGGVLYAQQWLSAGTHNGRDYALFFRQPV
      190     200     210     220     230     240

      250     260     270     280     290     300
    45 orf64ng-1.pep PENVAQDAVLIEKARAKYAE LSYSKKGLQTFFLVTLIIASLLSIFLALVMALYFARRFVE
       orf64-1      PKGVAEDAVLIEKARAKYAE LSYSKKGLQTFFLATLLIASLLSIFLALVMALYFARRFVE
      250     260     270     280     290     300

      310     320     330     340     350     360
    50 orf64ng-1.pep PILSLAEGAKAVAQGDFSQTRPYLRNDEFGRLTKLFNHMTQLSIAKEADERNRRREEAA
       orf64-1      PVLSLAEGAKAVAQGDFSQTRPYLRNDEFGRLTKLFNHMTQLSIAKEADERNRRREEAA
      310     320     330     340     350     360

      370     380     390     400     410     420
    55 orf64ng-1.pep RHYLECVLDGLTTGVVVVFDEKGRKTFNKAAEQILGMPLAPLWGSSRHGW HGVSAQQSLL
       orf64-1      RHYLECVLEGLTTGVVVVFDEQGCCLKTFNKAAEQILGMPLTPPLWGSSRHGW HGVSAQQSLL
      370     380     390     400     410     420

      430     440     450     460     470     480
    60 orf64ng-1.pep AEVFAAIGAAAGTDKPVQVEYAAPDDAKILLGKATVLPEDNGNGVVMVIDDITVLIRAQK
       orf64-1      AEVFAAIGAAAGTDKPVHVKYAAPDDAKILLGKATVLPEDNGNGVVMVIDDITVLIHAQK
      430     440     450     460     470     480

      490     500     510     520     530     540
    65 orf64ng-1.pep EAAWGEVAKRLAHEIRNPLTPIQLSAERLAWKLGGKLLDDQDAQILTRSTDITTIKQVAALK
       orf64-1      EAAWGEVAKRLAHEIRNPLTPIQLSAERLAWKLGGKLLDDQDAQILTRSTDITTIKQVAALK
      490     500     510     520     530     540
  
```


5	orf64-1	EEAWGEVAKRLAHEIRNPLTPTIQLSAERLAWKLGKLEQDAQILTRSTDTIVKQVAALK	490	500	510	520	530	540
	orf64ng-1.pep	EMVEAFERNYARAPSLKLENQDLNALIGDVLALYEAGPCRFEAEELAGEPLMMAADTTAMRQ	550	560	570	580	590	600
10	orf64-1	EMVEAFERNYARAPSLKLENQDLNALIGDVLALYEAGPCRFAAEELAGEPLTVAADTTAMRQ	550	560	570	580	590	600
	orf64ng-1.pep	VLHNIFFKNAAEEAADMPEVRVKSETGQDGRIVLTVCDNGKGFGEMLHNAFEPYVTDK	610	620	630	640	650	660
15	orf64-1	VLHNIFFKNAAEEAADMPEVRVKSETGQDGRIVLTVCDNGKGFGEMLHNAFEPYVTDK	610	620	630	640	650	660
	orf64ng-1.pep	PAGTGLGLPVVKIIEHGGRISSNQDAGGACVRILPKTVETYAX	670	680	690	700		
20	orf64-1	PAGTGLGLPVVKIIEHGGRISSNQDAGGACVRILPKTVETYAX	670	680	690	700		

Furthermore, ORF64ng-1 shows significant homology to a protein from *A. caulinodans*:

25	sp Q04850 NTRY_AZOCA NITROGEN REGULATION PROTEIN NTRY >gi 77479 pir S18624 ntry protein - Azorhizobium caulinodans >gi 38737 (X63841) NtrY gene product [Azorhizobium caulinodans] Length = 771 Score = 218 bits (550), Expect = 7e-56 Identities = 195/720 (27%), Positives = 320/720 (44%), Gaps = 58/720 (8%)							
	Query: 7	IAAICAVVLLYGLTAATGSTSSLDYFWWIXXXXXXXXXXXXXXXXXXRYVILLKDRRNGV	66					
30	Sbjct: 35	ISALATFLILMGLTPVVPVTHQVVIS---VLLVNAAVLILSAMVGREIWRIRAKARAGR	90					
	Query: 67	FGSQIAKRLSGMFTLVAVLPGLFLFGISAQFINGTINSWFGNDTHEALERSLNLKSALD	126					
35	Sbjct: 91	AAARLHIRIVGLFAVVSVPAILVAVVASLTDRGLDRWFSMRTQEIVASSVSVAQTYVR	150					
	Query: 127	LAADNAVSNAPVQIDLIGTASLSGNMGSVLEHYAG--SGFAQLALYNAASGKIEKSINP	184					
40	Sbjct: 151	EHALNIRGDILAMSADLTRLSKV-----YEGDRSRFNQILTAQAALRNLPGAMLI	200					
	Query: 185	HQFDQPLPKHEHWEIQQTGSVRSLESIGGVLYAQGWLSAGTHNGRDYA-----	233					
45	Sbjct: 201	RR-DLSVVERAN-VNIGREFIVPANLAIGDATPDQPVIIYLP--NDADYVAAVVPLKDYDD	256					
	Query: 234	--LFFRQPIPENVAQDAVLIEKARAKYAELSYSKKGLQTFFLVTXXXXXXXXXXXXVMA	291					
50	Sbjct: 257	LYLYVARLIDPRVIGYLKTTQETLADYRSLEERRFGVQVAFALMYAVITLIVLLSAVWL	316					
	Query: 292	LYFARRFVEPILSLAEGAKAVAQGFDSQTRPVLRLND-EFGRITKLFNMTQELSIXXXXX	350					
55	Sbjct: 317	LNFSKWLVAPIRRLMSAADHVAEGNLDVRVPIYRAEGDLASLAETFNKMTHELRSQREAI	376					
	Query: 351	XXXXXXXXXXHYLECVLDGLTTGVVVFDEKGRKLTFNKAAEQILGMPLAPLWGSSRHGW	410					
60	Sbjct: 377	LTARDQIDSRRRFTEAVLSGVGAGVIGLDSQERITILNRSARLLG--LSEVEALHRLA	434					
	Query: 411	HGVSAQQSLLAEVFXXXXXXXXTDKPVQVEYAAPDDAKILLGKATVLPEDNG---NGVVM	467					
65	Sbjct: 435	EVVPETAGLLEA-----EHARQSVQGNITLTRDGRERVFVRVTTEQSPEAEHGWV	488					
	Query: 468	VIDDITVLIRAQKEAAWGEVAKRLAHEIRNPLTPTIQLSAERLAWKLGKLEDDQDAQILTR	527					
70	Sbjct: 489	TLDDITELISAQRTSAWADVARRIAHEIKNPLTPTIQLSAERLKRKFGRHV-TQDREIFDQ	547					
	Query: 528	STDITIKQVAALKEMVEAFERNYARAPSLKLENQDLNALIGDVLALYEAGPCRFEAEELAGE	587					
75	Sbjct: 548	CTDTIIRQVGDIGRMVDFSSFARMKPVVDSQDMSEIIRQTVFLMRVGHPEVVDSEVP	607					
	Query: 588	PLMMAA-DTTAMRQVLHNIFFKNXXXXXXXXXDMPEVRVK-----SETGQDGRIVLTVCD	639					
80	Sbjct: 608	PAMPARFDRRLVSQALTNILKNAAEAIEAVP-PDVRGQGRIRVSANRVGED--LVIDIID	664					

Query: 640 NGKGFGEKMLHNAFEPPYVTDKPA GTGLGLPVVKIIEHGGGRISLSNQDAG-GACVRIIL 698
 NG G +E + EPYVT + GTGLGL +V KI+ EHGG I L++ G GA +R+ L
 Sbjct: 665 NGTGLPQESRNLLEPYVTTREKGTGLGLAIVGKIMEEHGGGIELNDAPEGRGAWIRLTL 724

Based on this analysis, including the presence of a putative leader sequence (double-underlined) and several putative transmembrane domains (single-underlined) in the gonococcal protein, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 31

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 259>:

```

      1 ATGTACGCAT TTACCGCCGC ACAGCAACAG AAGGCACTCT TCCGGCTGGT
    51 GCTTTTTCAT ATCCTCATCA TCGCCGCCAG CAACTATCTG GTGCAGTTCC
   101 CTTTCCAAAT TTTCGGCATC CACACCACTT GGGGCGCATT TTCCTTTCCC
   151 TTCATCTTCC TTGCCACCGA CCTGACCGTC CGCATTTTCG GTTCTCACTT
   201 GGCACGGCGG ATTATCTTTT GGGTGATGTT CCCCGCCCTT TTGCTTTCCT
   251 ACGTCTTTTC CGTTTGTTC CACAACGGCA GTTGGACAGG CTTGGGCGCG
   301 CTGTCCGAAT TCAACACCTT TGTCCGACGC ATCGCCTTAG CCAGCTTTGC
   351 CGCCTACGCG ATCGGACAAA TCCTTGATAT TTTTGTATTG AACAAATTAC
   401 GCCGTCTGAA AGCGTGGTGG ATTGCACCGA ACGCATCAAC CGTCATCGGG
   451 CACGCGTTGG ATACG...
```

This corresponds to the amino acid sequence <SEQ ID 260; ORF66>:

```

      1 MYAFTAAQQQ KALFRLVLFH ILIIAASNYL VQFPFQIFGI HTTWGAESFP
    51 FIFLATDLTV RIFGSHLARR IIFWVMFPAL LLSYVFSVLF HNGSWTGLGA
   101 LSEFNTFVGR IALASFAAYA IGQILDIFVF NKLRLKAWW IAPNASTVIG
   151 HALDT...
```

Further work revealed the complete nucleotide sequence <SEQ ID 261>:

```

      1 ATGTACGCAT TTACCGCCGC ACAGCAACAG AAGGCACTCT TCCGGCTGGT
    51 GCTTTTTCAT ATCCTCATCA TCGCCGCCAG CAACTATCTG GTGCAGTTCC
   101 CTTTCCAAAT TTTCGGCATC CACACCACTT GGGGCGCATT TTCCTTTCCC
   151 TTCATCTTCC TTGCCACCGA CCTGACCGTC CGCATTTTCG GTTCTCACTT
   201 GGCACGGCGG ATTATCTTTT GGGTGATGTT CCCCGCCCTT TTGCTTTCCT
   251 ACGTCTTTTC CGTTTGTTC CACAACGGCA GTTGGACAGG CTTGGGCGCG
   301 CTGTCCGAAT TCAACACCTT TGTCCGACGC ATCGCCTTAG CCAGCTTTGC
   351 CGCCTACGCG ATCGGACAAA TCCTTGATAT TTTTGTATTG AACAAATTAC
   401 GCCGTCTGAA AGCGTGGTGG ATTGCACCGA CCGCATCAAC CGTCATCGGC
   451 AACGCCTTGG ATACGCTGGT ATTTTTCGCC GTTGCCTTCT ACGCAAGCAG
   501 CGATGGATTT ATGGCGGCAA ACTGGCAGGG CATCGCTTTT GTCGATTACC
   551 TGTTCAAAC TACCGTCTGC ACCCTCTTCT TCCTGCCCGC CTACGGCGTG
   601 ATACTGAATC TGCTGACGAA AAAACTGACA ACCCTGCAAA CCAAACAGGC
   651 GCAAGACCGC CCCGCGCCCT CGCTGCAAAA TCCGTAA
```

This corresponds to the amino acid sequence <SEQ ID 262; ORF66-1>:

```

      1 MYAFTAAQQQ KALFRLVLFH ILIIAASNYL VQFPFQIFGI HTTWGAESFP
    51 FIFLATDLTV RIFGSHLARR IIFWVMFPAL LLSYVFSVLF HNGSWTGLGA
   101 LSEFNTFVGR IALASFAAYA IGQILDIFVF NKLRLKAWW IAPTASTVIG
   151 NALDTLVFFA VAFYASSDGF MAANWQGIAF VDYLFKLTVC TLFFLPAYGV
   201 ILNLLTKKLT TLQTKQAQDR PAPSLQNP*
```

Computer analysis of this amino acid sequence gave the following results:

Homology with the hypothetical protein o221 of *E. coli* (accession number P37619)

ORF66 and o221 protein show 67% aa identity in 155aa overlap:

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```

orf66 1 MYAFTAAQQQKALFRLVLFHILIIAASNYLVQFPFQIFGIHTTWGAFSFPFIFLATDLTV 60
M F+ Q+ KALF L LFH+L+I +SNYLQ P I G HTTWGAFSFPFIFLATDLTV
o221 1 MNVFSQTQRYKALEWLSLFHLLVITSSNYLVQLPVSILGFHTTWGAFSFPFIFLATDLTV 60

5 orf66 61 RIFGSHLARRIIFWVMFPALLLSYVFSVLFHNGSWTGLGALSEFNTFVGRIALASFAAYA 120
RIFG+ LARRIIF VM PALL+SYV S LF+ GSW G GAL+ FN FV RIA ASF AYA
o221 61 RIFGAPLARRIIFAVMIPALLISYVISSLFYMGSWQGFGLAHFNLFVARIATASFMAAYA 120

10 orf66 121 IGQILDIFVFNKLRLKAWWIAPNASTVIGHALDT 155
+GQILD+ VFN+LR+ + WW+AP AST+ G+ DT
o221 121 LGQILDVHVFENRLRQSRWWLAPTASTLFGNVSDT 155

```

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF66 shows 96.1% identity over a 155aa overlap with an ORF (ORF66a) from strain A of *N.*

15 *meningitidis*:

```

                10      20      30      40      50      60
orf66.pep      MYAFTAAQQQKALFRLVLFHILIIAASNYLVQFPFQIFGIHTTWGAFSFPFIFLATDLTV
                |||||
orf66a          MYAFTAAQQQKALEWLSLFHILIIAASNYLVQFPFQISGIHTTWGAFSFPFIFLATDLTV
                10      20      30      40      50      60

                70      80      90      100     110     120
orf66.pep      RIFGSHLARRIIFWVMFPALLLSYVFSVLFHNGSWTGLGALSEFNTFVGRIALASFAAYA
                |||||
25 orf66a          RIFGSHLARRIIFWVMFPALLLSYVFSVLFHNGSWTGLGALSEFNTFVGRIALASFAAYA
                70      80      90      100     110     120

                130     140     150
orf66.pep      IGQILDIFVFNKLRLKAWWIAPNASTVIGHALDT
                :|||
30 orf66a          LGQILDIFVFNKLRLKAWWVAPTASTVIGNALDTLVFFAVAFYASSDGFMAANWQGIAF
                130     140     150     160     170     180

orf66a          VDYLFKLTVCGLFFLPAYGVILNLLTKKLTTLQTKQAQDRPAPSLQNPX
                190     200     210     220

```

The complete length ORF66a nucleotide sequence <SEQ ID 263> is:

```

1  ATGTACGCAT TTACCGCCGC ACAGCAACAG AAGGCACTCT TCTGGCTGGT
51  GCTTTTTCAT ATCCTCATCA TCGCCGCCAG CAACTATCTG GTGCAGTTCC
40 101  CCTTCCAAAT TTCCGGCATC CACACCACTT GGGCGCGGTT TTCCTTTCCC
151  TTCATCTTCC TCGCCACCGA CCTGACCGTC CGCATTTTCG GTTCGCACCT
201  GGCACGGCGG ATTATCTTTT GGGTCATGTT CCGCGCCCTT TTGCTTTCCT
251  ACGTCTTTTC CGTTTGTTC CACAACGGCA GTTGGACGGG CTTGGGCGCG
301  CTGTCCGAAT TCAACACCTT TGTCGGACGC ATCGCGCTGG CAAGTTTTCG
351  CGCCTACGCG CTCGGACAAA TCCTTGATAT TTTTGTGTTC AACAAATTAC
45 401  GCCGTCTGAA AGCGTGGTGG GTTGCCCGA CTGCATCAAC CGTCATCGGC
451  AACGCCTTAG ATACGTTGGT ATTTTTCGCC GTTGCCCTCT ACGCAAGCAG
501  CGATGGATTT ATGGCGGCAA ACTGGCAGGG CATCGCTTTT GTCGATTACC
551  TGTTCAAACCT CACCGTCTGC GGTCTGTTT TCCTGCCCCG CTACGGCGTG
601  ATTCTGAATC TGCTGACGAA AAAACTGACG ACCCTGCAAA CCAACAGGC
50 651  GCAAGACCGC CCCGCGCCCT CGCTGCAAAA TCCGTAA

```

This encodes a protein having amino acid sequence <SEQ ID 264>:

```

1  MYAFTAAQQQ KALEWLVLFH ILIIAASNYL VQFPFQISGI HTTWGAFSFP
51  FIFLATDLTV RIFGSHLARR IIFWVMFPAL LLSYVFSVLF HNGSWTGLGA
101  LSEFNTFVGR IALASFAAYA LGQILDIFVF NKLRLKAWW VAPTASTVIG
55 151  NALDTLVFFA VAFYASSDGF MAANWQGIAF VDYLFKLTVC GLFFLPAYGV
201  ILNLLTKKLT TLQTKQAQDR PAPSLQNP*

```

ORF66a and ORF66-1 show 97.8% identity in 228 aa overlap:

```

                10      20      30      40      50      60
orf66a.pep      MYAFTAAQQQKALEWLVLFHILIIAASNYLVQFPFQISGIHTTWGAFSFPFIFLATDLTV
                |||||
60 orf66-1          MYAFTAAQQQKALFRLVLFHILIIAASNYLVQFPFQIFGIHTTWGAFSFPFIFLATDLTV

```

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		10	20	30	40	50	60
		70	80	90	100	110	120
5	orf66a.pep	RIFGSHLARRIIFWVMFPALLLSYVFSVLFHNGSWTGLGALSEFNTFVGRIALASFAAYA					
	orf66-1	RIFGSHLARRIIFWVMFPALLLSYVFSVLFHNGSWTGLGALSEFNTFVGRIALASFAAYA					
		70	80	90	100	110	120
10	orf66a.pep	LGQILDIFVFNKLRLKAWVAPTASTVIGNALDTLVFFAVAFYASSDGFMAANWQGI AF					
	orf66-1	IGQILDIFVFNKLRLKAWIAPTASTVIGNALDTLVFFAVAFYASSDGFMAANWQGI AF					
		130	140	150	160	170	180
15	orf66a.pep	VDYLFKLTVCGLFFLPAYGVILNLLTKKLTTLQTKQAQDRPAPSLQNPX					
	orf66-1	VDYLFKLTVCGLFFLPAYGVILNLLTKKLTTLQTKQAQDRPAPSLQNPX					
		190	200	210	220	229	
20							

Homology with a predicted ORF from *N.gonorrhoeae*

ORF66 shows 94.2% identity over a 155aa overlap with a predicted ORF (ORF66.ng) from *N.*

gonorrhoeae:

25	orf66.pep	MYAFTAAQQQKALFRLVLFHILIIAASNYLVQFPFQIFGIHTTWGAFSFPFIFLATDLTV	60
	orf66ng	MYALTAQQQKALFRLVLFHILIIAASNYLVQFPFRIFGIHTTWGAFSFPFIFLATDLTV	60
	orf66.pep	RIFGSHLARRIIFWVMFPALLLSYVFSVLFHNGSWTGLGALSEFNTFVGRIALASFAAYA	120
30	orf66ng	RIFGSHLARRIIFWVMFPALSLSYVFSVLFHNGSWTGLGAPSQFNTFVGRIALASFAAYA	120
	orf66.pep	IGQILDIFVFNKLRLKAWWIAPNASTVIGHALDT	155
	orf66ng	LGQILDIFVFDKLRLKAWWIAPAASTVIGNALDTLVFFAVAFYASSDEFMAANWQGI AF	180

35 The complete length ORF66ng nucleotide sequence <SEQ ID 265> is:

```

1  ATGTACGCAT TGACCGCCGC ACAGCAACAG AAGGCACTCT TCCGGCTGGT
51  GCTTTTCCAT ATCCTCATCA TCGCCGCCAG CAACTATCTG GTGCAGTTCC
101 CCTTCCGGAT TTTCGGCATC CACACCACTT GGGGCGCGTT TTCCTTTCCC
151 TTCATCTTCC TCGCCACCGA CCTGACCGTC CGCATTTTCG GTTCGCACTT
201 GCGCGCGCGG ATTATCTTTT GGGTGATGTT CCCC GCCCTT ttgCTTTCat
251 aCGTCTTTTC CGTTTGTTC CACAACGGCA GTTGGACGGG CTGGGCGCG
301 ctgTCCCAAT TCAACACCTT TGTCCGACGC ATCGCGCTGG CAAGTTTTC
351 CGCCTACGCG CTCGGACAAA TCCTTGATAT TTTCGTATTC GACAAATTAC
401 GCCGTCTGAA AGCGTGGTGG ATTGCCCGCG CCGCATCAAC CGTCATCGGC
451 AATGCACTGG ACACGTTAGT ATTTTGTGCC GTTGCTTTT ACGCAAGCAG
501 CGATGAATTT ATGGCGGCAA ACTGGCAGGG CATCGCTTT GTCGATTACC
551 TGTTCAAAC TACCGTCTGC ACCCTCTTCT TCCTGCCCGC CTACGGCGTG
601 ATACTGAATC TGCTGACGAA AAAACTGACG GCCCTGCAAA CCAAACAGGC
651 GCAAGACCGC CCCGTGCCCT CGCTGCAAAA TCCGTAA

```

50 This encodes a protein having amino acid sequence <SEQ ID 266>:

```

1  MYALTAQQQ KALFRLVLFH ILIIAASNYL VQFPFRIFGI HTTWGAFSFP
51  FIFLATDLTV RIFGSHLARR IIFWVMFPAL LSYVFSVLH HNGSWTGLGA
101 PSQFNTFVGR IALASFAAYA LGQILDIFVF DKLRLKAWW IAPAASTVIG
151 NALDTLVFFA VAFYASSDEF MAANWQGI AF VDYLFKLTVC TLFFLPAYGV
201 ILNLLTKKLT ALQTKQAQDR PVP SLQNP*

```

An alternative annotated sequence is:

```

1  MYALTAQQQ KALFRLVLFH ILIIAASNYL VQFPFRIFGI HTTWGAFSFP
51  FIFLATDLTV RIFGSHLARR IIFWVMFPAL LSYVFSVLH HNGSWTGLGA
101 LSQFNTFVGR IALASFAAYA LGQILDIFVF DKLRLKAWW IAPAASTVIG
151 NALDTLVFFA VAFYASSDEF MAANWQGI AF VDYLFKLTVC TLFFLPAYGV
201 ILNLLTKKLT ALQTKQAQDR PVP SLQNP*

```

ORF66ng and ORF66-1 show 96.1% identity in 228 aa overlap:

```

5      orf66-1.pep  MYAFTAAQQQKALFRLVLFHILIIAASNYLVQFPFQIFGIHTTWGAFSFPFIFLATDLTV  60
      orf66ng      MYALTAQQQKALFRLVLFHILIIAASNYLVQFPFRIFGIHTTWGAFSFPFIFLATDLTV  60
      orf66-1.pep  RIFGSHLARRIIFWVMFPALLLSYVFSVLFHNGSWTGLGALSEFNTEVGRIALASFAAYA 120
      orf66ng      RIFGSHLARRIIFWVMFPALLLSYVFSVLFHNGSWTGLGALSQFNTEVGRIALASFAAYA 120
10     orf66-1.pep  IGQILDIFVFNKLRRLKAWWIAPTASTVIGNALDTLVFFAVAFYASSDGFMAANWQGI AF 180
      orf66ng      LGQILDIFVFDKLRRLKAWWIAPAASTVIGNALDTLVFFAVAFYASSDEFMAANWQGI AF 180
      orf66-1.pep  VDYLEFKLTVCTLFFLPAYGVILNLLTKKLTTLQTKQAQDRPAPSLQNPX  229
15     orf66ng      VDYLEFKLTVCTLFFLPAYGVILNLLTKKLTALQTKQAQDRPVPSLQNPX  229

```

Furthermore, ORF66ng shows significant homology with an *E.coli* ORF:

```

20     sp|P37619|YHHQ_ECOLI HYPOTHETICAL 25.3 KD PROTEIN IN FTSY-NIKA INTERGENIC
      REGION (O221)
      >gi|1073495|pir||S47690 hypothetical protein o221 - Escherichia coli >gi|466607
      (U00039) No definition line found [Escherichia coli] >gi|1789882 (AE000423)
      hypothetical 25.3 kD protein in ftsY-nikA intergenic region [Escherichia coli]
      Length = 221
25     Score = 273 bits (692), Expect = 5e-73
      Identities = 132/203 (65%), Positives = 155/203 (76%)

      Query: 1  MYALTAQQQKALFRLVLFHILIIAASNYLVQFPFRIFGIHTTWGAFSFPFIFLATDLTV  60
      M + Q+ KALF L LFH+L+I +SNYLVQ P I G HTTWGAFSFPFIFLATDLTV
      Sbjct: 1  MNVFSQTQRYKALFWLSLFHLLVITSSNYLVQLPVSI LGFHTTWGAFSFPFIFLATDLTV  60
30     Query: 61 RIFGSHLARRIIFWVMFPALLLSYVFSVLFHNGSWTGLGALSQFNTEVGRIALASFAAYA 120
      RIFG+ LARRIIF VM PALL+SYV S LF+ GSW G GAL+ FN FV RIA ASF AYA
      Sbjct: 61 RIFGAPLARRIIFAVMIPALLISYVISSLFYMGSWQGF GALAHFNLFVARIATASEMAYA 120
35     Query: 121 LGQILDIFVFDKLRRLKAWWIAPAASTVIGNALDTLVFFAVAFYASSDEFMAANWQGI AF 180
      LGQILD+ VF++LR+ + WW+AP AST+ GN DTL FF +AF+ S D FMA +W IA
      Sbjct: 121 LGQILDVHVFNRLRQSRWWLAPTASTLFGNVSDTLAFFFI AFWRSPDAEAEHWMEIAL 180
40     Query: 181 VDYLEFKLTVCTLFFLPAYGVILN 203
      VDY FK+ + +FFLP YGV+LN
      Sbjct: 181 VDYCFKVLISIVFFLPMYGVLLN 203

```

Based on this analysis, including the homology with the *E.coli* protein and the presence of several putative transmembrane domains in the gonococcal protein, it is predicted that these proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 32

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 267>:

```

50     1  ATGGTCATAA AATATACAAA TTTGAATTTT GCGAAATTGT CGATAATTGC
      51  AATTTTGATG ATGTATTCGT TTGAAGCGAA TGCAAAyGCA GTmwrAATAT
      101 CTGAAACTGT TTCAGTTGAT ACCGGACAAG GTGCGAAAAT TCATAAGTTT
      151 GTACCTAAAA ATAGTAAAAC TTATTCATCT GATTTAATAA AAACGGTAGA
      201 TTAAACACAC AyyCCTACGG GCGCAAAAGC CCGAATCAAC GCCAAAATAA
      251 CCGCCAGCGT ATCCCGCGCC GCGGTATTGG CGGGGGTCGG CAAACTTGCC
55     301 CGCTTAGgCG CGAAATTCAG CACAAGGGCG GTtCCCTATG TCGGAACAGC
      351 CcTTTTAGCC CACGACGTAT ACGAAAcTTT CAAAGAAGAC ATACAGGCAC
      401 GAGGCTACCA ATACAGCCCC GAAACCGACA AATTTGTAAA AGGCTACGAA
      451 TATAGTAATT GCCTTTGGTA CGAAGACAAA AGACGTATTA ATAGAACCTA

```

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501 TGGCTGCTAC GCGTTGAT..

This corresponds to the amino acid sequence <SEQ ID 268; ORF72>:

5 1 MVIKYTNLNF AKLSIIAILM MYSFEANANA VXISETVSVD TGQGAKIHKF
 51 VPKNSKTYSS DLIKTVDLTH XPTGAKARIN AKITASVSRA GVLAVGVKLA
 101 RLGAKFSTRA VPYVGTALLA HDVYETFKED IQARGYQYDP ETDKFVKGYE
 151 YSNCLWYEDK RRINRTYGCY GVD..

Further work revealed the complete nucleotide sequence <SEQ ID 269>:

10 1 ATGGTCATAA AATATACAAA TTTGAATTTT GCGAAATTGT CGATAATTGC
 51 AATTTTGATG ATGTATTCGT TTGAAGCGAA TGCAAATGCA GTAAAAATAT
 101 CTGAAACTGT TTCAGTTGAT ACCGGACAAG GTGCGAAAAT TCATAAGTTT
 151 GTACCTAAAA ATAGTAAAAC TTATTCATCT GATTTAATAA AAACGGTAGA
 201 TTTAACACAC ATCCCTACGG GCGCAAAAGC CCGAATCAAC GCCAAAATAA
 251 CCGCCAGCGT ATCCCGCGCC GCGTATTGG CGGGGGTCGG CAAACTTGCC
 301 CGCTTAGGCG CGAAATTCAG CACAAGGGCG GTTCCCTATG TCGGAACAGC
 15 351 CCTTTAGCC CACGACGTAT ACGAACTTT CAAAGAAGAC ATACAGGCAC
 401 GAGGCTACCA ATACGACCCC GAAACCGACA AATTGCAAA GGTCTCAGGC
 451 TAA

This corresponds to the amino acid sequence <SEQ ID 270; ORF72-1>:

20 1 MVIKYTNLNF AKLSIIAILM MYSFEANANA VKISETVSVD TGQGAKIHKF
 51 VPKNSKTYSS DLIKTVDLTH IPTGAKARIN AKITASVSRA GVLAVGVKLA
 101 RLGAKFSTRA VPYVGTALLA HDVYETFKED IQARGYQYDP ETDKFAKVS
 151 *

Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

25 ORF72 shows 98.0% identity over a 147aa overlap with an ORF (ORF72a) from strain A of *N. meningitidis*:

		10	20	30	40	50	60
orf72.pep		MVIKYTNLNF AKLSIIAILM MYSFEANANA VXISETVSVD TGQGAKIHKE VPKNSKTYSS					
30 orf72a		MVIKYTNLNF AKLSIIAILM MYSFEANANA VKISETVSVD TGQGAKIHKE VPKNSKTYSS					
		10	20	30	40	50	60
		70	80	90	100	110	120
orf72.pep		DLIKTVDLTH XPTGAKARIN AKITASVSRA GVLAVGVKLARLGA KFSTRAPVYVGTALLA					
35 orf72a		DLIKTVDLTH IPTGAKARIN AKITASVSRA GVLAVGVKLARLGA KFSTRAPVYVGTALLA					
		70	80	90	100	110	120
		130	140	150	160	170	
40 orf72.pep		HDVYETFKEDI QARGYQYDP ETDKFVKGYE YSNCLWYEDK RRINRTYGCY GVD					
orf72a		HDVYETFKEDI QARGYQYDP ETDKFAKVS GX					
		130	140	150			

The complete length ORF72a nucleotide sequence <SEQ ID 271> is:

45 1 ATGGTCATAA AATATACAAA TTTGAATTTT GCGAAATTGT CGATAATTGC
 51 AATTTTGATG ATGTATTCGT TTGAAGCGAA TGCAAATGCA GTAAAAATAT
 101 CTGAAACTGT TTCAGTTGAT ACCGGACAAG GTGCGAAAAT TCATAAGTTT
 151 GTACCTAAAA ATAGTAAAAC TTATTCATCT GATTTAATAA AAACGGTAGA
 201 TTTAACACAC ATCCCTACGG GCGCAAAAGC CCGAATCAAC GCCAAAATAA
 50 251 CCGCCAGCGT ATCCCGCGCC GCGTATTGG CGGGGGTCGG CAAACTTGCC
 301 CGCTTAGGCG CGAAATTCAG CACAAGGGCG GTTCCCTATG TCGGAACAGC
 351 CCTTTAGCC CACGACGTAT ACGAACTTT CAAAGAAGAC ATACAGGCAC
 401 GAGGCTACCA ATACGACCCC GAAACCGACA AATTGCAAA GGTCTCAGGC
 451 TAA

55 This encodes a protein having amino acid sequence <SEQ ID 272>:

-193-

```

1  MVIKYTNLNF AKLSIIAILM MYSFEANANA VKISETVSVD TGQGAKIHKF
51 VPKNSKTYSS DLIKTVDLTH IPTGAKARIN AKITASVSRA GVLAVGVGKLA
101 RLGAKFSTRA VPYVGTALLA HDVYETFKED IQARGYQYDP ETDKFAKVS
151 *

```

5 ORF72a and ORF72-1 show 100.0% identity in 150 aa overlap:

```

10 orf72a.pep      10      20      30      40      50      60
    MVIKYTNLNF AKLSIIAILM MYSFEANANA VKISETVSVD TGQGAKIHKF VPKNSKTYSS
    |||
10 orf72-1        10      20      30      40      50      60
    MVIKYTNLNF AKLSIIAILM MYSFEANANA VKISETVSVD TGQGAKIHKF VPKNSKTYSS
    |||

15 orf72a.pep      70      80      90     100     110     120
    DLIKTVDLTH IPTGAKARIN AKITASVSRA GVLAVGVGKLARLGAKFSTRAVPYVGTALLA
    |||
15 orf72-1        70      80      90     100     110     120
    DLIKTVDLTH IPTGAKARIN AKITASVSRA GVLAVGVGKLARLGAKFSTRAVPYVGTALLA
    |||

20 orf72a.pep      130     140     150
    HDVYETFKEDI QARGYQYDP ETDKFAKVS GX
    |||
20 orf72-1        130     140     150
    HDVYETFKEDI QARGYQYDP ETDKFAKVS GX
    |||

```

Homology with a predicted ORF from *N.gonorrhoeae*

25 ORF72 shows 89% identity over a 173aa overlap with a predicted ORF (ORF72.ng) from *N. gonorrhoeae*:

```

30 orf72.pep      MVIKYTNLNF AKLSIIAILM MYSFEANANA VKISETVSVD TGQGAKIHKF VPKNSKTYSS 60
    |||
30 orf72ng        MVTKHTNLNF AKLSIIAILM MYSFEANANA VKISETLSVD TGQGAKVHKF VPKSSNIYSS 60
    |||

35 orf72.pep      DLIKTVDLTH XPTGAKARIN AKITASVSRA GVLAVGVGKLARLGAKFSTRAVPYVGTALLA 120
    |||
35 orf72ng        DLTKAVDLTH IPTGAKARIN AKITASVSRA GVLAVGVGKLVRQGAKFSTRAVPYVGTALLA 120
    |||

35 orf72.pep      HDVYETFKEDI QARGYQYDP ETDKFKVKG YEYSNCLWYED KRRINR TYGCGYVD 173
    |||
35 orf72ng        HDVYETFKEDI QARGCRYDP ETDKFKVKG YEYANCLWYED DERRINR TYGCGYVD SSIMRLM 180
    |||

```

An ORF72ng nucleotide sequence <SEQ ID 273> was predicted to encode a protein having amino acid sequence <SEQ ID 274>:

```

40 1  MVTKHTNLNF AKLSIIAILM MYSFEANANA VKISETLSVD TGQGAKVHKF
51 VPKSSNIYSS DLTKAVDLTH IPTGAKARIN AKITASVSRA GVLAVGVGKLV
101 RQGAKFGTRA VPYVGTALLA HDVYETFKED IQARGCRYDP ETDKFKVKG YE
151 YANCLWYEDE RRINR TYGCGY GVDSSIMRLM PDRSRFPEVK QLMESQMYRL
201 ARPFWNRKE ELNKLSSLDW NNFVLNRCTF DWNGGGCAVN KGDDFRAGAS
45 251 FSLGRNPKYK EEMDAKKPEE ILSLKVDADP DKYIEATGYP GYSEKVEVAP
301 GTKVNMGPVT DRNGNPVQVA ATFGRDAQGN TTADVQVIPR PDLTPASAEA
351 PHAQPLPEVS PAENPANNDP DENPGTRPN PEPDPLNDP ANPDTGQPG
401 TSPDSPAVPD RPNGRHRKER KEGEDGGLSC DYFPEILACQ EMGKPSDRMF
451 HDISIPQVTD DKTWSSHNF LPSNGVCPQPK TFHVFGRRQYR ASYEPLCVFA
50 501 EKIRFAVLLA FIIMSAFVVF GSLGGE*

```

After further analysis, the following gonococcal DNA sequence <SEQ ID 275> was identified:

```

55 1  ATGGTCACAA AACATACAAA TTTGAATTTT GCGAAATTGT CGATAATTGC
51 AATTTTGATG ATGTATTCGT TTGAAGCGAA TGCAATGCA GTAAAAATAT
101 CTGAAACTCT TTCGGTTGAT ACCGACAAAG GCGCGAAAGT TCATAAGTTC
151 GTTCCTAAAT CAAGTAATAT TTATTCATCT GATTTAACAA AAGCGGTAGA
201 TTTAACGCAT ATCCCCACGG GCGCAAAAGC CCGAATCAAC GCCAAATATA
251 CCGCCAGCGT ATCCCCGCGC GCGGTATTGT CGGGGGTCGG CAAACTTGTC
301 CGCCAAGCGC CGAAATTCGG CACAAGGGCG GTTCCCTATG TCGGAACAGC
351 CCTTTTAGCC CACGACGTAT ACGAACTTT CAAAGAAGAC ATACAGGCAC
60 401 GAGGCTGCCG ATACGATCCC GAAACCGACA AATTT

```

This corresponds to the amino acid sequence <SEQ ID 276; ORF72ng-1>:

```

1  MVTKHTNINF AKLSIIAILM MYSFEANANA VKISETLSVD TGQGAQVHKF
51  VPKSSNIYSS DLTKAVDLTH IPTCAKARIN AKITASVSRA GVLSGVGKLV
101 RQGAQFGTRA VEVYGTALLA HDVYETFKED IQARGCRYDP ETDKF

```

5 ORF72ng-1 and ORF721-1 show 89.7% identity in 145 aa overlap:

```

10 orf72ng-1.pe 10 20 30 40 50 60
    MVTKHTNINF AKLSIIAILM MYSFEANANA VKISETLSVD TGQGAQVHKF VPKSSNIYSS
    || :|||||
orf72-1 10 20 30 40 50 60
    MVIKYTNLNF AKLSIIAILM MYSFEANANA VKISETVSDT GQGAQVHKF VPKNSKTYSS

15 orf72ng-1.pe 70 80 90 100 110 120
    DLTKAVDLTH IPTGAKARIN AKITASVSRA GVLSGVGKLV RQGAQFGTRA VYVGTALLA
    || :|||||
orf72-1 15 70 80 90 100 110 120
    DLIKTVDLTH IPTGAKARIN AKITASVSRA GVLGAGVGLAR LGAKFSTRAPVYVGTALLA

20 orf72ng-1.pe 130 140
    HDVYETFKEDI QARGCRYDP ETDKF
    ||| :|||
orf72-1 20 130 140 150
    HDVYETFKEDI QARGYQYDP ETDKFAKVS GX

```

Based on this analysis, including the presence of a putative leader sequence and transmembrane domains in the gonococcal protein, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 33

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 277>:

```

30 1  ATGAGATTTT TCGGTATCGG TTTTGTGGTG CTGCTGTTTT TGGAGATTAT
51  GTCGATTGTG TGGGTTGCCG ATTGGCTGGG CGGCGGCTGG ACGTTGTTTT
101 TGATGGCGGC AGGTTTGGCC GCCGGCGTGC TGATGCTCAG GCAAACCGGG
151 CTGACCCGGT CTTTATTGGG CGGCGCGGCG AATGAGAAGC GGCGGGAAGG
201 TATCCGTTTA TCAGATGTTG TGGCCTATC..

```

35 This corresponds to the amino acid sequence <SEQ ID 278; ORF73>:

```

1  MRFFGIGFLV LLFLEIMSIV WVADWLGGGW TLFLMAAGFA AGVLMRLRQTG
51  LTGLLLAGAA MRSGGKVS VY QMLWPI..

```

Further work revealed the complete nucleotide sequence <SEQ ID 279>:

```

40 1  ATGAGATTTT TCGGTATCGG TTTTGTGGTG CTGCTGTTTT TGGAGATTAT
51  GTCGATTGTG TGGGTTGCCG ATTGGCTGGG CGGCGGCTGG ACGTTGTTTT
101 TGATGGCGGC AGGTTTGGCC GCCGGCGTGC TGATGCTCAG GCATACGGGC
151 CTGTCCGGTC TTTTATTGGC GGGCGCGGCA ATGAGAAGCG GCGGGAGGGT
201 ATCCGTTTAT CAGATGTTGT GGCCTATCCG TTATACGGTG GCGGCTGTGT
251 GTCTGATGAG TCCGGGATTC GTATCCTCGG TGTTGGCGGT ATTGCTGCTG
45 301 CTGCCGTTTA AGGGAGGGGC AGTGTTCAG GCAGGAGGTG CGGAAAATTT
351 TTCAACATG AACCAATCGG GCAGAAAAGA GGGCTTTTCC CGCGATGACG
401 ATATTATCGA GGGAGAATAT ACGGTTGAAG AGCCTTACGG CGGCAATCGT
451 TCCCGAAACG CCATCGAACA CAAAAAGAC GAATAA

```

This corresponds to the amino acid sequence <SEQ ID 280; ORF73-1>:

```

50 1  MRFFGIGFLV LLFLEIMSIV WVADWLGGGW TLFLMAAGFA AGVLMRLRHTG
51  LSGLLLAGAA MRSGGRVSVY QMLWPIRYTV AAVCLMSPGF VSSVLAVLLL
101 LPFKGGAVLQ AGGAENFFNM NQSGRKEGFS RDDDIIEGEY TVEEPYGGNR

```


151 SRNAIEHKKD E*

Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF73 shows 90.8% identity over a 76aa overlap with an ORF (ORF73a) from strain A of *N.*

5 *meningitidis*:

```

      10      20      30      40      50      60
orf73.pep MRFFGIGFLVLLFLEIMSIWVADWLGGGWTFLFLMAAGFAAGVLMRLRQTGLTGLLLAGAA
          |||||:|||||
orf73a    MRFFGIGFLVLLFLEIMSIWVADWLGGGWTFLFLMAATFAAGVVMRLRHTGLSGLLLAGAA
      10      20      30      40      50      60

      70
orf73.pep MRSGGKVSQMLWPI
          |||||:|||||
orf73a    MRSGGRVSVYXMLWXIRYTVAAVCXMSPGFVSSVXAVLLXLPFKGGAVLQAGGAENFFNM
15

```

The complete length ORF73a nucleotide sequence <SEQ ID 281> is:

```

      1  ATGAGATTTT TCGGTATCGG TTTTGTGGTG CTGCTGTTTT TGGAGATTAT
     51  GTCGATTGTG TGGGTTGCCG ATTGGTTGGG CGGCGGTTGG ACGCTGTTTC
    101  TAATGGCGGC AACCTTTGCC GCCGGCGTGG TGATGCTCAG GCATACGGGG
    151  CTGTCCGGTC TTTTATTTGGC GGGCGCGGCA ATGAGAAGCG GCGGGAGGGT
    201  ATCCGTTTAT CANATGTTGT GGCNTATCCG TTATACGGTG GCGGCGGTGT
    251  GTCNGATGAG TCCGGGATTC GTATCCTCGG TGTNGGCGGT ATTGCTGNTG
    301  CTNCCGTTTA AGGGAGGTGC AGTGTTGCAG GCAGGAGGTG CGGAAAATTT
    351  TTTCAACATG AACCANTCGG GCAGAAAAGA NGGCNTTTCC CGCGATGACG
    401  ATATTATCGA GGGGGAATAT ACGGTTGAAG ANCCTTACGG CGGCANTCGT
    451  TTCCGAAACG CCNTNGAACA CAAAAAAGAC GAATAA

```

This encodes a protein having amino acid sequence <SEQ ID 282>:

```

      1  MRFFGIGFLV LLFLEIMSIW VVADWLGGGW TLFLMAATFA AGVVMRLRHTG
     51  LSGLLLAGAA MRSGGRVSVY XMLWXIRYTV AAVCXMSPGF VSSVXAVLLX
    101  LPFKGGAVLQ AGGAENFFNM NXSGRKXGXS RDDDIIEGEY TVEXPYGGXR
    151  FRNAXEHKKD E*

```

ORF73a and ORF73-1 show 91.3% identity in 161 aa overlap

```

      10      20      30      40      50      60
orf73a.pep MRFFGIGFLVLLFLEIMSIWVADWLGGGWTFLFLMAATFAAGVVMRLRHTGLSGLLLAGAA
          |||||:|||||
orf73-1    MRFFGIGFLVLLFLEIMSIWVADWLGGGWTFLFLMAAGFAAGVLMRLRHTGLSGLLLAGAA
      10      20      30      40      50      60

      70      80      90     100     110     120
orf73a.pep MRSGGRVSVYXMLWXIRYTVAAVCXMSPGFVSSVXAVLLXLPFKGGAVLQAGGAENFFNM
          |||||:|||||
orf73-1    MRSGGRVSVYQMLWPIRYTVAAVCLMSPGFVSSVLAVLLLPFKGGAVLQAGGAENFFNM
      70      80      90     100     110     120

      130     140     150     160
orf73a.pep NXSGRKXGXS RDDDIIEGEYTVEXPYGGXRFRNAXEHKKDEX
          |||||:|||||
orf73-1    NQSGRKEGFSRDDDIIEGEYVEEPYGGNRSRNAIEHKKDEX
      130     140     150     160

```

Homology with a predicted ORF from *N.gonorrhoeae*

ORF73 shows 92.1% identity over a 76aa overlap with a predicted ORF (ORF73.ng) from *N.*

gonorrhoeae:

```

      10      20      30      40      50      60
orf73.pep MRFFGIGFLVLLFLEIMSIWVADWLGGGWTFLFLMAAGFAAGVLMRLRQTGLTGLLLAGAA
          |||||:|||||

```

```

      orf73ng      MRFFGIGFLVLLFLEIMSIVWVADWLGCGWTFLMAATFAAGVLMRLHTGLSGLLLAGAA      60
      orf73.pep    MRSGBKVSQYQLWPI                                                    76
                    ::||:|||||
5      orf73ng      VKSSGKVSQYQLWPIRYTVAAVCLMSPGFVSSVLAVLLLLLPPKGGAVLQAGGAENFFNM    120

```

The complete length ORF73ng nucleotide sequence <SEQ ID 283> is:

```

      1  ATGAGATTTT TCGGTATCGG TTTTGTGGTG CTGCTGTTTT TGGAAATTAT
      51  GTCGATTGTG TGGGTGCGCG ATTGGCTGGG CGGCGGTTGG AcgcTGTTC
     101  TAATGGCGGC AACCTTTGCC GCCGGTGTGC TGATGCTCAG GCATAcggGG
     151  CTGTCCGGTC TTTTATTGGC TGGCGCGGCG GTAAAAagta gtgGGAAGGT
     201  ATCTGTTTAT CagatgtTGT GGCTATCCG TTATAcggtg gcggcggtgT
     251  GTCTGatgag tCcgGATTC GTATCCTccg tgttgCGGT ATTGCTGCTG
     301  CTGCcgttta aggGaggGgc agtggtgcag gcaggaggtg cggaaaATTT
     351  TTTCAACATg aaCcaatcgg gcagaaAaga gggatttttc cacgatgacg
     401  atattatcga gggagaatat acggttgaaa aacctgacgg cggcaatcgt
     451  tcccgaAAcG ccatcgaaca cgaaaAgac gaataA

```

This encodes a protein having amino acid sequence <SEQ ID 284>:

```

      1  MRFFGIGFLV LLFLEIMSIV WVADWLGGGW TLFLMAATFA AGVLMRLHTG
     51  LSGLLLAGAA VKSSGKVSQY QMLWPIRYTV AAVCLMSPGF VSSVLAVLLL
     101  LPPKGGAVLQ AGGAENFFNM NQSGRKEGFF HDDDIIEGEY TVEKPDGGRN
     151  SRNAIEHEKD E*

```

ORF73ng and ORG73-1 show 93.8% identity in 161 aa overlap

```

      10      20      30      40      50      60
25  orf73-1.pep  MRFFGIGFLVLLFLEIMSIVWVADWLGCGWTFLMAAGFAAGVLMRLHTGLSGLLLAGAA
      orf73ng    MRFFGIGFLVLLFLEIMSIVWVADWLGCGWTFLMAATFAAGVLMRLHTGLSGLLLAGAA
      10      20      30      40      50      60
30  orf73-1.pep  MRSGBRVSVYQLWPIRYTVAAVCLMSPGFVSSVLAVLLLLLPPKGGAVLQAGGAENFFNM
      orf73ng    VKSSGKVSQYQLWPIRYTVAAVCLMSPGFVSSVLAVLLLLLPPKGGAVLQAGGAENFFNM
      70      80      90      100     110     120
35  orf73-1.pep  :||:|||||
      orf73ng    VKSSGKVSQYQLWPIRYTVAAVCLMSPGFVSSVLAVLLLLLPPKGGAVLQAGGAENFFNM
      70      80      90      100     110     120
40  orf73-1.pep  NQSGRKEGFSRDDDIIEGEYTVPEPYGGNRSRNAIEHKKDEX
      orf73ng    NQSGRKEGFFHDDDIIEGEYTVKPDGGRNRSRNAIEHEKDEX
      130     140     150     160

```

Based on this analysis, including the presence of a putative leader sequence and putative transmembrane domain in the gonococcal protein, it is predicted that the proteins from *N.meningitidis* and *N.gonorrhoeae*, and their epitopes, could be useful antigens for vaccines or diagnostics, or for raising antibodies.

Example 34

The following partial DNA sequence was identified in *N.meningitidis* <SEQ ID 285>:

```

      1  ATGTTTGT TTACAGACGGC ATTCTT.ATG TTTCAGAAAC ATTTGCAGAA
      51  AGCCTCCGAC AGCGTCGTCG GAGGGACATT ATACGTGGTT GCCACGCCCA
     101  TCGCAATTT GCGGACATT ACCGTGCGCG CTTGGCGGT ATTGCAAAAG
     151  GCG..... .GCCGA AGACACGCGC GTTACCGCAC AGCTTTTGAG
     201  CGCGTACGGC ATTCAGGGCA AACTCGTCAG TGTGCGCGAA CACAACGAAC
     251  GGCAGATGGC GGACAAGATT GTCGGCTATC TTTCAGACGG CATGGTTGTG
     301  GCACAGGTTT CCGATGCGGG TACGCCGGCC GTGTGCGACC CGGGCGCGAA
     351  ACTCGCCCGC CGCGTGCCTG AGGCCGGGTT TAAAGTCGTT CCCGTCGTGG
     401  GCGCAAC.GC GGTGATGGCG GCTTTGAGCG TGGCCGGTGT GGAAGGATCC
     451  GATTTTATT TCAACGGTTT TGTACCGCGC AAATCGGGAG AACGCAGGAA

```

501 ACTGTTTGCC AAATGGGTGC GGGCGGCGTT TCCTATCGTC ATGTTTGAAA
 551 CGCCGCACCG CATCGGTGCA GCGCTTGCCG ATATGGCGGA ACTGTTCCCC
 601 GAACGCCGAT TAATGCTGGC GCGCGAAATT ACGAAAACGT TTGAAACGTT
 651 CTTAAGCGGC ACGGTTGGGG AAATTCAGAC GGCATTGTCT GCCGACGGCG
 701 ACCAATCGCG CGGCGAGATG GTGTTGGTGC TTTATCCGCG GCAGGATGAA
 751 AAACACGAAG GCTTGTCCGA GTCCGCGCAA AACATCATGA AAATCCTCAC
 801 AGCCGAGCTG CCGACCAAAC AGGCGGCGGA GCTTGCTGCC AAAATCACGG
 851 GCGAGGGAAA GAAAGCTTTG TACGAT..

This corresponds to the amino acid sequence <SEQ ID 286; ORF75>:

1 MFVFQTAFXM FQKHLQKASD SVVGGTLYV V ATPIGNLADI TLRALAVLQK
 51 A...AEDTR VTAQLLSAYG IQKLVSVRE HNERQMADKI VGYLSDGMVV
 101 AQVSDAGTPA VCDPGAKLAR RVREAGFKVV PVVGAXAVMA ALSVAGVEGS
 151 DFYENGFPVP KSGERRKLFA KVVRAAFPIV MFETPHRIGA ALADMAELFP
 201 ERRMLLAREI TKTFETFLSG TVGEIQTALS ADGDQSRGEM VLVLYPAQDE
 251 KHEGLSESAQ NIMKILTAE L PTKQAELAA KITGEGKKAL YD..

Further work revealed the complete nucleotide sequence <SEQ ID 287>:

1 ATGTTTCAGA AACATTTGCA GAAAGCCTCC GACAGCGTCG TCGGAGGGAC
 51 ATTATACGTG GTTGCCACGC CCATCGGCAA TTTGGCGGAC ATTACCCTGC
 101 GCGCTTTGGC GGTATTGCAA AAGGCGGACA TCATCTGTGC CGAAGACACG
 151 CCGCTTACCG CACAGCTTTT GAGCGCGTAC GGCATTCAGG GCAAACCTCGT
 201 CAGTGTGCGC GAACACAACG AACGCGAGAT GCGGACAAAG ATTGTCGGCT
 251 ATCTTTCAGA CGGCATGGTT GTGGCACAGG TTTCCGATGC GGGTACGCCG
 301 GCCGTGTGCG ACCCGGGCGC GAAACTCGCC CGCGCGTGC GTGAGGCCGG
 351 GTTTAAAGTC GTTCCCCTCG TGGGCGCAAG CGCGGTGATG GCGGCTTTGA
 401 GCGTGGCCGG TGTGGAAGGA TCCGATTTT ATTCAACG TTTGTACCG
 451 CCGAAATCGG GAGAACGCG GAAACTGTTT GCCAAATGGG TCGGGGCGGC
 501 GTTTCCTATC GTCATGTTT GAAACGCGCA CCGCATCGGT GCGACGCTTG
 551 CCGATATGGC GGAAGTGTTC CCCGAACGCC GATTAATGCT GGCAGCGCAA
 601 ATTACGAAAA CGTTTGAAAC GTTCTTAAGC GGCACGGTTG GGGAAATCA
 651 GACGGCATTG TCTGCCGACG GCAACCAATC GCGCGGCGAG ATGGTGTGG
 701 TGCTTTATCC GCGCGAGGAT GAAAAACACG AAGGCTTGTC CGAGTCCGCG
 751 CAAAACATCA TGAATTCCT CACAGCCGAG CTGCCGACCA AACAGGCGGC
 801 GAGAGCTTGT GCCAAATCA CGGCGGAGG AAAGAAAGT TGTACGATC
 851 TGGCTCTGTC TTGAAAAAC AAATAG

35 This corresponds to the amino acid sequence <SEQ ID 288; ORF75-1>:

1 MFQKHLQKAS DSVVGGTLYV VATPIGNLAD ITLRALAVLQ KADIICAEDT
 51 RVTAQLLSAY GIQKLVSVR EHNERQMAK I VGYLSDGMV VAQVSDAGTP
 101 AVCDPGAKLA RRVREAGFKV VPVVGASAVM AALSVAGVEG SDFYFNGFVP
 151 PKSGERRKLF AKWVRAAFPI VMFETPHRIG ATLADMAELF PERRMLLARE
 201 ITKTFETFLS GTVGEIQTAL SADGNQSRGE MVLVLYPAQD EKHEGLSESA
 251 QNIMKILTAE LPTKQAELA AKITGEGKKA LYDLALSWKN K*

Computer analysis of this amino acid sequence gave the following results:

Homology with a predicted ORF from *N.meningitidis* (strain A)

ORF75 shows 95.8% identity over a 283aa overlap with an ORF (ORF75a) from strain A of *N.*

45 *meningitidis*:

		10	20	30	40	50	60
orf75.pep		MFVFQTAFXM	FQKHLQKASD	SVVGGTLYV	VATPIGNLAD	ITLRALAVLQ	KAXXXAEDTR
orf75a		MFQKHLQKASD	SVVGGTLYV	VATPIGNLAD	ITLRALAVLQ	KADIICAEDTR	
50		10	20	30	40	50	
		70	80	90	100	110	120
orf75.pep		VTAQLLSAYGI	QKLVSVREHNER	QMAKIVGYLSDGMV	VAQVSDAGT	PAVCDPGAKLAR	
orf75a		VTAQLLSAYGI	QKLVSVREHNER	QMAKIVGYLSDGMV	VAQVSDAGT	PAVCDPGAKLAR	
55		60	70	80	90	100	110
		130	140	150	160	170	180
orf75.pep		RVREAGFKV	VPVVGAXAVMA	ALSVAGVEGS	DFYFNGFVP	PKSGERRKL	FAKVVRAAFPIV

15

The complete length ORF75a nucleotide sequence <SEQ ID 289> is:

35

This encodes a protein having amino acid sequence <SEQ ID 290>:

40

ORF75a and ORF75-1 show 98.3% identity in 291 aa overlap:

65